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Type 2 diabetes care: the role of insulinsensitizing agents and practical implications for cardiovascular disease prevention

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Available online 15 November 1998.

Abstract

Millions of Americans are at risk for cardiovascular morbidity and mortality related to disorders of glucose intolerance—particularly type 2 diabetes and prediabetic conditions, including the insulin resistance, or "cardiovascular dysmetabolic," syndrome. The latter is apparently more intricately associated with macrovascular disease—myocardial infarction, stroke, and peripheral vascular disease. In some situations the risk of cardiovascular disease might be reduced by the prevention of diabetes and also by prevention or treatment of the cardiovascular dysmetabolic syndrome. Studies have shown that intensive glycemic control can delay the development of microvascular complications in type 1, and possibly type 2, diabetes. Several longitudinal observational studies have demonstrated a relationship between glycemic control and the development of cardiovascular disease. Prospective clinical intervention trials to address this issue are underway. Insulin may have a role in atherogenesis, both directly and by promoting development of such risk factors as hypertension and dyslipidemia. Genetic factors and mechanisms promoting or discouraging development of glucose intolerance are also under investigation. Lifestyle changes—dietary and exercise modification, weight loss, and smoking cessation—have been shown to have a positive effect on cardiovascular disease risk. Clinical trials suggest that oral antidiabetic agents—particularly the new noninsulin secretagogues (including troglitazone and

metformin, which act on the liver and on skeletal muscle)—may be useful in delaying or preventing development of type 2 diabetes and the cardiovascular dysmetabolic syndrome, as well as in their treatment, when present. Both agents, acting primarily by different mechanisms of action, have also demonstrated potential beneficial effects on serum lipid profiles and other cardiovascular risk factors and may be useful in patients with cardiovascular dysmetabolic syndrome who do not yet meet the criteria for diabetes.

Requests for reprints should be addressed to Robert R. Henry, MD, Veterans Administration Hospital, 3350 La Jolla Village Drive, La Jolla, California 92161

The American Journal of Medicine Volume 105, Issue 1, Supplement 1, 6 July 1998, Pages 20S-26S This Document

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L3 439 S THIAZOLIDINE 2 4 DIONE

9 S PYRIDYL AMINO ETHOXY BENZYL

L5 1 S L4 AND L3

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```
=> s 15 or rosiglitazone
  19 FILES SEARCHED...
  32 FILES SEARCHED...
          9168 L5 OR ROSIGLITAZONE
L6
=> s rosiglitazone maleate
           603 ROSIGLITAZONE MALEATE
=> s diabet?
  28 FILES SEARCHED...
       1608844 DIABET?
L8
=> s type II
  20 FILES SEARCHED...
        414754 TYPE II
=> s 18 (s) 19
  29 FILES SEARCHED...
         42916 L8 (S) L9
L10
=> s 17 and 110
           119 L7 AND L10
L11
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=> d l12 100-115 ibib, kwic
L12 ANSWER 100 OF 115 COPYRIGHT 2003 Gale Group on STN
                    1998:153994 NLDB
ACCESSION NUMBER:
                    Rosiglitazone Success In Phase III Boosts SB
TITLE:
                    Marketletter, (22 Jun 1998) .
SOURCE:
                    ISSN: 0951-3175.
                    Marketletter Publications Ltd. (UK)
PUBLISHER:
DOCUMENT TYPE:
                    Newsletter
                    English
LANGUAGE:
                     609
WORD COUNT:
     SmithKline Beecham's insulin sensitizer Avandia (rosiglitazone
     maleate) has been shown to significantly reduce blood glucose
      levels in patients with type II diabetes,
      according to data from a Phase III clinical trial presented at the recent
      American Diabetes Association annual meeting held in Chicago,
      USA.
                  . . recently when the US National Institutes of Health
      discontinued the troglitazone arm of its five-year trial for the
      prevention of type II diabetes after one
      patient developed liver failure and subsequently died following
      transplantation (Marketletter June 15).
                                      COPYRIGHT 2003 IMSWORLD on STN
L12 ANSWER 101 OF 115 DRUGLAUNCH
      Trade Name: AVANDAMET
 CN
       Chemical Name: metformin hydrochloride; rosiglitazone
 CN
       maleate; metformin hydrochloride; rosiglitazone
       maleate; metformin hydrochloride; rosiglitazone
       maleate
 COMP Active Ingredient: tabs film-coated a: metformin hydrochloride, 500 mg,
                          rosiglitazone maleate, 1 mg; tabs
                          film-coated b: metformin hydrochloride, 500 mg,
                          rosiglitazone maleate, 2 mg; tabs
```

film-coated c: metformin hydrochloride, 500 mg,

rosiglitazone maleate, 4 mg.

- TX Type II diabetes.
- L12 ANSWER 102 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate
- COMP Active Ingredient: rosiglitazone maleate, 4 mg.
- TX Combined oral therapy of type II diabetes
 - mellitus in patients insufficiently controlled with metformin or sulfonylurea derivatives.
- L12 ANSWER 103 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate;

rosiglitazone maleate

COMP Active Ingredient: tabs film-coated a: rosiglitazone maleate, 4 mg; tabs film-coated b:

rosiglitazone maleate, 4 mg.

- TX Treatment of type II diabetes
- L12 ANSWER 104 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate
- COMP Active Ingredient: rosiglitazone maleate, 4 mg.
- TX Treatment of diabetes mellitus (type II)
- L12 ANSWER 105 OF 115 DRUGLAUNCH COPYRIGHT 2003, IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate
- COMP Active Ingredient: rosiglitazone maleate, 4 mg.
- TX Diabetes mellitis type II
- L12 ANSWER 106 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate
- COMP Active Ingredient: rosiglitazone maleate, 4 mg.
- TX Treatment of type II diabetes mellitus
- L12 ANSWER 107 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate;

rosiglitazone maleate

COMP Active Ingredient: tabs film-coated a: rosiglitazone maleate, 4 mg; tabs film-coated b:

rosiglitazone maleate, 4 mg.

- TX Treatment of type II diabetes
- L12 ANSWER 108 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA
- CN Chemical Name: rosiglitazone maleate
- COMP Active Ingredient: rosiglitazone maleate, 4 mg.
- TX Treatment of type II diabetes
- L12 ANSWER 109 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN
- CN Trade Name: AVANDIA

maleate

- CN Chemical Name: rosiglitazone maleate; rosiglitazone maleate; rosiglitazone
- COMP Active Ingredient: tabs film-coated a: rosiglitazone maleate, 4 mg; tabs film-coated b: rosiglitazone maleate, 4 mg; tabs

film-coated c: rosiglitazone maleate, 8 mg.

TX . . . combination with metformin in obese patients, or with a sulphonylurea in patients with an intolerance or contraindication to metformin, in type II diabetes inadequately controlled by maximal tolerated doses of either.

L12 ANSWER 110 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN

CN Trade Name: AVANDIA

CN Chemical Name: rosiglitazone maleate;

rosiglitazone maleate

COMP Active Ingredient: tabs coated a: rosiglitazone maleate , 4 mg; tabs coated b: rosiglitazone maleate, 8 mg.

TX Treatment of type II diabetes mellitus

L12 ANSWER 111 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN

CN Trade Name: AVANDIA

CN Chemical Name: rosiglitazone maleate

COMP Active Ingredient: rosiglitazone maleate, 4 mg.

TX Treatment of type II diabetes

L12 ANSWER 112 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN

CN Trade Name: AVANDIA

CN Chemical Name: rosiglitazone maleate; rosiglitazone maleate; rosiglitazone maleate

COMP Active Ingredient: tabs film-coated a: rosiglitazone maleate, 2 mg; tabs film-coated b: rosiglitazone maleate, 4 mg; tabs film-coated c: rosiglitazone maleate

, 8 mg.

TX Diabetes mellitus type II

L12 ANSWER 113 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN

CN Trade Name: AVANDIA

CN Chemical Name: rosiglitazone maleate; rosiglitazone maleate; rosiglitazone maleate; rosiglitazone maleate; rosiglitazone maleate; rosiglitazone maleate

COMP Active Ingredient: tabs coated a: rosiglitazone maleate
, 2 mg; tabs coated b: rosiglitazone
maleate, 2 mg; tabs coated c:
rosiglitazone maleate, 4 mg; tabs
coated d: rosiglitazone maleate, 4
mg; tabs coated e: rosiglitazone
maleate, 8 mg; tabs coated f:
rosiglitazone maleate, 8 mg.

TX Treatment of type II diabetes

L12 ANSWER 114 OF 115 DRUGLAUNCH COPYRIGHT 2003 IMSWORLD on STN

CN Trade Name: AVANDIA

CN Chemical Name: rosiglitazone maleate; rosiglitazone maleate; rosiglitazone maleate

COMP Active Ingredient: tabs a: rosiglitazone maleate, 2
mg; tabs b: rosiglitazone maleate,
4 mg; tabs c: rosiglitazone maleate
, 8 mg.

TX Treatment of type II diabetes

L12 ANSWER 115 OF 115 PHARMAML COPYRIGHT 2003 MARKETLETTER ON STN ACCESSION NUMBER: 1642110 PHARMAML

Rosiglitazone Success In Phase III Boosts SB TITLE:

Marketletter June 18, 1998 SOURCE:

Newsletter DOCUMENT TYPE:

599 WORD COUNT:

SmithKline Beecham's insulin sensitizer Avandia (rosiglitazone maleate) has been shown to significantly reduce blood glucose levels in patients with type II diabetes,

according to data from a Phase III clinical trial presented at the recent American Diabetes Association annual meeting held in Chicago,

USA.

. recently when the US National Institutes of Health discontinued the troglitazone arm of its five-year trial for the prevention of type II diabetes after one patient developed liver failure and subsequently died following transplantation

=> d 112 90-99 ibib, kwic

L12 ANSWER 90 OF 115 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN

DUPLICATE 4

ACCESSION NUMBER: 2000:106894 BIOSIS PREV200000106894 DOCUMENT NUMBER:

(Marketletter June 15).

Hepatocellular injury in a patient receiving rosiglitazone: TITLE:

A case report.

Al-Salman, Jameela; Arjomand, Heider; Kemp, David G.; AUTHOR (S):

Mittal, Manoj (1)

(1) Division of Gastroenterology, Easton Hospital, 250 CORPORATE SOURCE:

South 21st Street, Easton, PA, 18042 USA

Annals of Internal Medicine, (Jan. 18, 2000) Vol. 132, No. SOURCE:

2, pp. 121-124.

ISSN: 0003-4819.

DOCUMENT TYPE: Article English LANGUAGE: SUMMARY LANGUAGE: English

Background: Rosiglitazone maleate (Avandia, SmithKline

Beecham, Philadelphia, Pennsylvania) is a new oral hypoglycemic agent

approved for the treatment of type 2 diabetes. It. . .

IT Major Concepts

> Gastroenterology (Human Medicine, Medical Sciences); Pharmacology; Toxicology

IT

hepatocellular injury: digestive system disease; type II diabetes: endocrine disease/pancreas, metabolic disease

IT Chemicals & Biochemicals

rosiglitazone maleate: antidiabetic - drug

Alternate Indexing

Diabetes Mellitus, Non-Insulin-Dependent (MeSH)

155141-29-0 (ROSIGLITAZONE MALEATE) RN

L12 ANSWER 91 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER: 1999:1817 DRUGNL

rosiglitazone SmithKline Beecham clinical data TITLE:

R&D Focus Drug News (28 Jun 1999). SOURCE:

WORD COUNT: 173

SmithKline Beecham reported clinical data on its therapy for noninsulin-dependent (type II) diabetes mellitus, rosiglitazone (AVANDIA) at the 81st Annual Meeting of the Endocrine Society, 15-21 June 1999, San Diego, USA. When administered. .

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 92 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER: 1999:1556 DRUGNL

TITLE: rosiglitazone Bristol-Myers Squibb, SmithKline Beecham

marketed, USA

SOURCE: R&D Focus Drug News (31 May 1999).

WORD COUNT: 84

TX The US FDA granted approval 26 May 1999 for SmithKline Beecham's thiazolidinedione rosiglitazone (AVANDIA) for use in type II diabetes. According to the company, the agent will be launched within a few days of approval. Rosiglitazone is an insulin sensitizer.

CN rosiglitazone; rosiglitazone maleate; BRL 49653; BRL 49653c; SB 210232; AVANDIA

CN rosiglitazone; rosiglitazone maleate; BRL 49653; BRL 49653c; SB 210232; AVANDIA

L12 ANSWER 93 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER: 1999:3206 DRUGNL

TITLE: rosiglitazone SmithKline Beecham rosiglitazone receives

negative review by EMEA

SOURCE: R&D Focus Drug News (1 Nov 1999).

WORD COUNT: 68

TX Rosiglitazone, indicated for type II diabetes
, has been marketed in the USA and Mexico and approved in a number of countries.

CN rosiglitazone; rosiglitazone maleate; BRL 49653; BRL
49653c; SB 210232; AVANDIA

L12 ANSWER 94 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER: 1999:818 DRUGNL

TITLE: rosiglitazone SmithKline Beecham awaits FDA committee

review

SOURCE: R&D Focus Drug News (29 Mar 1999).

WORD COUNT: 46

TX The . . . (AVANDIA), SmithKline Beecham's thiazolidinedione, in April 1999. The product had been submitted in the USA for the treatment of noninsulin-dependent (type II) diabetes mellitus. Approval is awaited in Europe.

CN rosiglitazone; rosiglitazone maleate; BRL 49653; BRL 49653c; SB 210232; AVANDIA

L12 ANSWER 95 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER: 1999:262 DRUGNL

TITLE: rosiglitazone SmithKline Beecham receives priority review

status

SOURCE: R&D Focus Drug News (1 Feb 1999).

WORD COUNT: 64

TX The US FDA has granted six-month priority review status to rosiglitazone (AVANDIA), SmithKline Beecham's insulin sensitizer, for the treatment of type II diabetes. The company submitted the NDA November 1998 seeking approval of rosiglitazone for use as a monotherapy or a combination therapy.. . .

CN rosiglitazone; rosiglitazone maleate; BRL 49653; BRL 49653c; SB 210232; AVANDIA

L12 ANSWER 96 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER: 1999:2184 DRUGNL

TITLE: rosiglitazone SmithKline Beecham marketed, Mexico

SOURCE: R&D Focus Drug News (26 Jul 1999).

WORD COUNT: 38

TX Rosiglitazone . . . sensitizer developed by SmithKline Beecham, has been launched in Mexico for use as monotherapy or in certain combination

regimens for type II diabetes. The product has been submitted for approval in 42 countries worldwide.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 97 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER:

1999:1171 DRUGNL

TITLE:

rosiglitazone SmithKline Beecham registered, Mexico SmithKline Beecham submitted for approval, Canada,

Australia SmithKline Beecham recommended for approval, USA

SmithKline Beecham licensing agreement Bristol-Myers

Squibb licensing agreement

SOURCE:

R&D Focus Drug News (3 May 1999).

WORD COUNT:

126

SmithKline . . . been recommended for approval by the US FDA's TX Endocrinologic and Metabolic Drugs Advisory Committee for the treatment of noninsulin dependent (type II) diabetes mellitus either as a monotherapy or in combination with metformin. In another development, SmithKline Beecham announced an agreement with Bristol-Myers.

The product has been approved for marketing in Mexico as a monotherapy and in certain combination regimens for treatment of type II diabetes. Approval of rosiglitazone is pending additionally in Europe, Australia and Canada.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 98 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER:

1998:4367 DRUGNL

TITLE:

rosiglitazone SmithKline Beecham submitted for approval,

Europe

SOURCE:

R&D Focus Drug News (14 Dec 1998).

WORD COUNT:

Closely . . . sensitizer, in the 15 countries of the European Union via the centralized procedure. The filing specifies rosiglitazone's use in treating type II (noninsulin-dependent) diabetes, either as a monotherapy or in combination with certain other agents.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 99 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

ACCESSION NUMBER:

1998:4292 DRUGNL

TITLE:

rosiglitazone SmithKline Beecham submitted for approval,

USA

SOURCE:

R&D Focus Drug News (7 Dec 1998).

WORD COUNT:

106

The . . . by developer SmithKline Beecham. The filing, the first TX worldwide for the product, is seeking approval for its use in treating type II (noninsulin-dependent) diabetes,

either as a monotherapy or in combination with certain other agents. Worldwide clinical trials of rosiglitazone are reported to have.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

=> d 112 80-89 ibib, kwic

L12 ANSWER 80 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:622 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham marketed, Central TITLE:

America, Colombia, Puerto Rico R&D Focus Drug News (21 Feb 2000).

26 WORD COUNT:

SOURCE:

. insulin sensitizer developed by SmithKline Beecham, Rosiglitazone . has been launched in Central America, Colombia and Puerto Rico for the treatment of type II diabetes.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 81 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:4103 DRUGNL ACCESSION NUMBER:

ramipril, rosiglitazone Aventis phase change III, TITLE:

Worldwide SmithKline Beecham

R&D Focus Drug News (11 Dec 2000). SOURCE:

103 WORD COUNT:

A . . . initiated to compare Aventis' ACE inhibitor ramipril (ALTACE) with SmithKline Beecham's thiazolidinedione insulin sensitizer rosiglitazone (AVANDIA) in the prevention of type II diabetes. The trial, known as DREAM (Diabetes REduction Approaches with ramipril and rosiglitazone Medications), aims to enroll 4000 subjects at high risk for developing type II diabetes owing to impaired glucose tolerance. Patients will receive placebo, ramipril, rosiglitazone or a combination of the two agents. The study.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 82 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:3275 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham marketed, Ireland, TITLE:

Finland

R&D Focus Drug News (16 Oct 2000). SOURCE:

21 WORD COUNT:

SmithKline Beecham's rosiglitazone (AVANDIA), an insulin sensitizer for the treatment of type II diabetes, has been launched in Ireland and Finland.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 83 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:772 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham, Asahi Chemical licensing TITLE:

agreement

R&D Focus Drug News (28 Feb 2000). SOURCE:

WORD COUNT:

SmithKline Beecham has signed a copromotion agreement with Asahi Chemical for its thiazolidinedione, rosiglitazone maleate (AVANDIA), in Japan. The compound, which is an insulin sensitizer, is a treatment for non-insulin dependent (type II) diabetes mellitus and has been marketed in a number of Latin

American countries and the USA, where a copromotion agreement exists. .

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 84 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:1082 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham recommended for approval, TITLE:

R&D Focus Drug News (27 Mar 2000). SOURCE:

WORD COUNT: 85

SmithKline Beecham's rosiglitazone (AVANDIA) has been recommended for approval by the Europe's CPMP for the treatment of noninsulin dependent (TX type II) diabetes mellitus in combination with other oral antidiabetic agents. This recommendation follows an appeal to the EMEA which had rejected rosiglitazone.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL 49653c; SB 210232; AVANDIA

L12 ANSWER 85 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:1215 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham marketed, Canada TITLE:

R&D Focus Drug News (10 Apr 2000). SOURCE:

WORD COUNT:

SmithKline . . . approval for and launched rosiglitazone (AVANDIA) in Canada for use alone or in combination with metformin in the treatment of type II diabetes. The compound, a thiazolidinedione insulin sensitizer, is already available in certain countries in North and South America and awaits approval. . .

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL 49653c; SB 210232; AVANDIA

L12 ANSWER 86 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:1115 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham marketed, Brazil TITLE:

R&D Focus Drug News (3 Apr 2000). SOURCE:

20 WORD COUNT:

Rosiglitazone (AVANDIA), an insulin sensitizer developed by SmithKline Beecham, has been launched Brazil for the treatment of type II diabetes.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 87 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:2792 DRUGNL ACCESSION NUMBER:

rosiglitazone SmithKline Beecham receives NICE TITLE:

recommendation

R&D Focus Drug News (4 Sep 2000). SOURCE:

WORD COUNT: SmithKline Beecham's once-daily antidiabetic, rosiglitazone (AVANDIA), has been recommended for use in combination with oral monotherapy for patients with type II diabetes by the National Institute for Clinical Excellence (NICE), UK. This advisory body on drug prescription stated that, in the light. .

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 88 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2000:2417 DRUGNL

ACCESSION NUMBER: rosiglitazone SmithKline Beecham registered, Europe TITLE:

R&D Focus Drug News (24 Jul 2000). SOURCE:

. by the European Commission for use in combination with WORD COUNT:

other oral antidiabetic drugs, under defined circumstances, in the

treatment of type II diabetes. SmithKline

Beecham is planning to commence product launches throughout the European Union, initially in the UK and Germany, during July.

Rosiglitazone has been approved for the treatment of type II diabetes in 38 countries worldwide, including the

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 89 OF 115 COPYRIGHT 2003 Gale Group on STN

2000:303951 NLDB ACCESSION NUMBER:

A compound discovered by ASAHI CHEMICAL INDUSTRY CO., TITLE: LTD. (Brief Article)

Japan-U.S. Business Report, (1 Mar 2000) pp. 2.

SOURCE: ISSN: 0888-5702.

Japan Economic Institute of America PUBLISHER:

Newsletter DOCUMENT TYPE: English LANGUAGE:

A compound discovered by ASAHI CHEMICAL INDUSTRY CO., LTD. that shows WORD COUNT:

promise in the treatment of obesity and diabetes will be exclusively developed, commercialized and sold outside East Asia by SMITHKLINE BEECHAM PLC. AZ40140 is a Beta-3 receptor agonist. . . and its partner will jointly commercialize AZ40140. As part of the agreement, Asahi Chemical will codistribute in Japan SB's Avandia (rosiglitazone maleate), a treatment for Type

II, or adult-onset, diabetes. That product is in Phase

II clinical testing there.

THIS IS THE FULL TEXT: COPYRIGHT 2000 Japan Economic Institute of.

=> d 112 1-10 ibib, kwic

L12 ANSWER 1 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2003:464 DRUGNL

ACCESSION NUMBER: rosiglitazone GlaxoSmithKline, Sankyo licensing agreement TITLE:

R&D Focus Drug News (10 Feb 2003). SOURCE:

65 WORD COUNT:

Rosiglitazone, a thiazolidinedione is marketed for the treatment of type II diabetes in many countries, including the USA, Canada and Europe. Approval is pending in Japan.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 2 OF 115 DRUGNL COPYRIGHT 2003 IMSWORLD on STN

2003:991 DRUGNL ACCESSION NUMBER:

rosiglitazone GlaxoSmithKline receives additional approval TITLE:

R&D Focus Drug News (17 Mar 2003). SOURCE:

WORD COUNT:

GlaxoSmithKline's thiazolidinedione, rosiglitazone (AVANDIA), has been approved by the US FDA for use in combination with insulin for the treatment of type II diabetes mellitus.

Consequently, the therapy is indicated as a monotherapy or as a combination therapy with metformin, sulfonylureas or insulin to improve glycemic control in patients with type II diabetes.

rosiglitazone; rosiglitazone maleate; BRL 49653; BRL CN 49653c; SB 210232; AVANDIA

L12 ANSWER 3 OF 115 USPATFULL on STN

ACCESSION NUMBER: 2003:258657 USPATFULL

TITLE:

Death domain-containing receptor polynucleotides,

polypeptides, and antibodies

INVENTOR (S):

Ni, Jian, Germantown, MD, UNITED STATES Ruben, Steven M., Olney, MD, UNITED STATES

Human Genome Sciences, Inc., Rockville, MD (U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE

PATENT INFORMATION: APPLICATION INFO.:

20030925 US 2003181710 **A1** A1 20020620 (10) US 2002-175042

RELATED APPLN. INFO.:

Continuation of Ser. No. US 2001-835788, filed on 17 Apr 2001, ABANDONED Continuation-in-part of Ser. No. WO

2000-US28666, filed on 17 Oct 2000, PENDING

NUMBER DATE

PRIORITY INFORMATION:

US 1999-159585P 19991018 (60) US 1999-167246P 19991124 (60)

DOCUMENT TYPE:

Utility APPLICATION

FILE SEGMENT: LEGAL REPRESENTATIVE:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS:

22 1

EXEMPLARY CLAIM: LINE COUNT:

14139

SUMM

. component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia, Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye.

SUMM

. . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, and/or ameliorate type 11 diabetes

mellitus (insulin resistant diabetes mellitus).

SUMM

. . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemic-hyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases".

. Papillary Adenocarcinoma SUMM H0684

Ovarian cancer, Serous Papillary Adenocarcinoma

Ovarian Cancer H0689

Ovarian Cancer, #9702G001

H0690 Prostate Adenocarcinoma H0696 Human Adult Skeletal Muscle

H0706 Adipose tissue (diabetic type I, obese) #41706 H0713

Adipose tissue (diabetic type II) H0717

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#41661
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Chromosome 22 exon L1290 Brain frontal cortex S0001 Monocyte activated S0002 Human Osteoclastoma S0003 Early Stage Human Brain S0007

disease

Human Amygdala

Stromal cell.

(glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM..

L12 ANSWER 4 OF 115

ACCESSION NUMBER:

TITLE:

S0010

S0026

DETD

INVENTOR (S):

USPATFULL on STN 2003:258639 USPATFULL

207 human secreted proteins

Ni, Jian, Germantown, MD, UNITED STATES Ebner, Reinhard, Gaithersburg, MD, UNITED STATES LaFleur, David W., Washington, DC, UNITED STATES Moore, Paul A., Germantown, MD, UNITED STATES Olsen, Henrik S., Gaithersburg, MD, UNITED STATES Rosen, Craig A., Laytonsville, MD, UNITED STATES Ruben, Steven M., Olney, MD, UNITED STATES Soppet, Daniel R., Centreville, VA, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Shi, Yanggu, Gaithersburg, MD, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES Wei, Ying-Fei, Berkeley, CA, UNITED STATES Florence, Charles, Rockville, MD, UNITED STATES Hu, Jing-Shan, Mountain View, CA, UNITED STATES Li, Yi, Sunnyvale, CA, UNITED STATES Kyaw, Hla, Frederick, MD, UNITED STATES Fischer, Carrie L., Burke, VA, UNITED STATES Ferrie, Ann M., Painted Post, NY, UNITED STATES Fan, Ping, Potomac, MD, UNITED STATES Feng, Ping, Gaithersburg, MD, UNITED STATES Endress, Gregory A., Florence, MA, UNITED STATES Dillon, Patrick J., Carlsbad, CA, UNITED STATES Carter, Kenneth C., North Potomac, MD, UNITED STATES Brewer, Laurie A., St. Paul, MN, UNITED STATES Yu, Guo-Liang, Berkeley, CA, UNITED STATES Zeng, Zhizhen, Lansdale, PA, UNITED STATES

NUMBER	KIND	DATE

PATENT INFORMATION: APPLICATION INFO .: RELATED APPLN. INFO.:

20030925 US 2003181692 A1 20010822 (9) **A1** US 2001-933767 Continuation-in-part of Ser. No. WO 2001-US5614, filed on 21 Feb 2001, PENDING Continuation-in-part of Ser. No. US 1998-205258, filed on 4 Dec 1998, PENDING

Greene, John M., Gaithersburg, MD, UNITED STATES

DATE

PRIORITY	INFORMATION:	US
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US	2000-184836P	20000224	(60)
	2000-193170P	20000329	(60)
US	1997-48885P	19970606	(60)
US	1997-49375P	19970606	(60)
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US	1997-48876P	19970606	(60)
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US	1997-48884P	19970606	(60)

NUMBER

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 US 1998-94657P
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Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, LEGAL REPRESENTATIVE: ROCKVILLE, MD, 20850 NUMBER OF CLAIMS: 23 EXEMPLARY CLAIM: 10 Drawing Page(s) NUMBER OF DRAWINGS: . component that may be treated, prevented, and/or diagnosed with LINE COUNT: the compositions of the invention include, but are not limited to, SUMM type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders. [2042] Moreover, disorders and/or states, which can be treated, prevented, diagnosed, and/or prognosed with the the polynucleotides, SUMM polypeptides, agonists and/or agonists of the invention include, but are not limited to , solid tumors, blood born tumors such as leukemias, tumor metastasis , Kaposi's sarcoma, benign tumors, for example hemangiomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas, rheumatoid arthritis, psoriasis, ocular angiogenic diseases, for example, diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis, retinoblastoma, and uvietis, delayed wound healing, endometriosis, vascluogenesis, granulations, hypertrophic scars (keloids), nonunion fractures, scleroderma, trachoma, vascular adhesions, myocardial angiogenesis, coronary collaterals, cerebral collaterals, arterioyenous malformations, ischemic limb angiogenesis, Osler -Webber Syndrome, plaque neovascularization, telangiectasia, hemophiliac joints, angiofibroma fibromuscular dysplasia, wound granulation, Crohn's disease, atherosclerosis, birth control agent by preventing vascularization required for embryo implantation controlling menstruation, diseases that have angiogenesis as a pathologic consequence such as cat scratch disease (Rochele minalia quintosa), ulcers (Helicobacter pylori), Bartonellosis and bacillary angiomatosis. (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. DETD (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTAT.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. [2657] The diabetic animals have many of the characteristic. DETD features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al., J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol. 83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67 (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D. L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous diabetic mice develop hyperglycemia that is resistant to insulin analogous to human type II diabetes

(Mandel et al., J. Immunol. 120:1375-1377 (1978)).

ACCESSION NUMBER:

2003:258441 USPATFULL

TITLE:

Novel heterocyclic analogs of diphenylethylene

compounds

INVENTOR(S):

Neogi, Partha, Fremont, CA, UNITED STATES

Dey, Debendranath, Fremont, CA, UNITED STATES

Medicherla, Satyanarayana, Cupertino, CA, UNITED STATES

Nag, Bishwajit, Union City, CA, UNITED STATES Lee, Arthur, San Francisco, CA, UNITED STATES

KIND DATE NUMBER -----

PATENT INFORMATION:

US 2003181494 A1 20030925 US 2002-265902 A1 20021008 (10)

APPLICATION INFO.: RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 2001-843167, filed on 27 Apr 2001, PENDING Continuation-in-part of Ser. No. US 2001-785554, filed on 20 Feb 2001, PENDING Continuation-in-part of Ser. No. US 2000-591105, filed on 9 Jun 2000, ABANDONED Continuation-in-part of Ser.

No. US 1999-287237, filed on 6 Apr 1999, GRANTED, Pat.

No. US 6331633

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

PILLSBURY WINTHROP, LLP, P.O. BOX 10500, MCLEAN, VA,

22102

NUMBER OF CLAIMS:

40

EXEMPLARY CLAIM:

1 26 Drawing Page(s)

NUMBER OF DRAWINGS:

2827

LINE COUNT: . which are effective in lowering blood glucose level, serum insulin, triglyceride and free fatty acid levels in animal models of

Type II diabetes. The compounds are

disclosed as useful for a variety of treatments including the treatment

of inflammation, inflammatory and immunological diseases,. . .

effects. For example, the compounds are useful in lowering blood glucose, serum insulin and triglyceride levels in animal models of SUMM

type II diabetes.

[0004] The causes of type I and type II SUMM

diabetes are yet unknown, although both genetics and environment seem to be major factors. Insulin dependent type I and non-insulin

dependent type II are the types which are known.

Type I is an autoimmune disease in which the responsible autoantigen is still unknown. Patients of type I need to take insulin parenterally or

subcutaneously to survive. However, type II

diabetes, the more common form, is a metabolic disorder

resulting from the body's inability to make a sufficient amount of

insulin. .

[0013] The thiazolidinedione class listed in the above table has gained SUMM more widespread use in recent years for treatment of type

II diabetes, exhibiting particular usefulness as

insulin sensitizers to combat "insulin resistance", a condition in which the patient becomes less responsive to.

. . . obtained from Jackson Laboratories (Bar Harbor, Me.) when their age was 5 weeks. Seven-week-old animals were dosed with Compound 11, rosiglitazone maleate (recrystallized from

commercially available tablets) or vehicle (0.5% carboxymethyl cellulose (Sigma, St. Louis, Mo.) in water) orally once daily by. . .

L12 ANSWER 6 OF 115 USPATFULL on STN

ACCESSION NUMBER:

2003:257883 USPATFULL

TITLE:

DETD

Plasminogen-like polynucleotides, polypeptides, and

antibodies

INVENTOR (S):

Ni, Jian, Germantown, MD, UNITED STATES

Young, Paul E., Gaithersburg, MD, UNITED STATES

Ruben, Steven M., Olney, MD, UNITED STATES Human Genome Sciences, Inc., Rockville, MD, UNITED

PATENT ASSIGNEE(S): STATES, 20850 (U.S. corporation)

> KIND NUMBER -----

US 2003180934 A1 20030925 US 2002-162742 A1 20020606 (10) PATENT INFORMATION: Continuation of Ser. No. US 2001-832197, filed on 11 APPLICATION INFO.: Apr 2001, ABANDONED Continuation-in-part of Ser. No. WO RELATED APPLN. INFO.: 2000-US27253, filed on 4 Oct 2000, PENDING DATE NUMBER -----US 1999-158044P 19991007 (60) PRIORITY INFORMATION: Utility DOCUMENT TYPE: APPLICATION HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, FILE SEGMENT: LEGAL REPRESENTATIVE: ROCKVILLE, MD, 20850 22 NUMBER OF CLAIMS: 1 EXEMPLARY CLAIM: . . . component that may be treated, prevented, and/or diagnosed with LINE COUNT: the compositions of the invention include, but are not limited to, SUMM type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia, Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye. . . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, SUMM and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus). . . . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated SUMM with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal

failure, nephropathy other diseases and disorders as described in the "Renal Discrders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". . . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.

(gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; . alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. .

L12 ANSWER 7 OF 115 USPATFULL on STN

DETD

2003:251897 USPATFULL ACCESSION NUMBER:

Retinoid receptor interacting polynucleotides, TITLE:

polypeptides, and antibodies

Shi, Yanggu, Gaithersburg, MD, UNITED STATES INVENTOR (S): Ruben, Steven M., Olney, MD, UNITED STATES

Human Genome Sciences, Inc., Rockville, MD, UNITED PATENT ASSIGNEE(S):

STATES (U.S. corporation)

DATE KIND NUMBER _____ US 2003176686 A1 20030918 US 2002-193159 A1 20020712 (10)

PATENT INFORMATION: APPLICATION INFO .: RELATED APPLN. INFO.:

Continuation of Ser. No. US 2001-788600, filed on 21 Feb 2001, ABANDONED Continuation-in-part of Ser. No. WO

2000-US22351, filed on 15 Aug 2000, PENDING

DATE NUMBER -----

19990816 (60) US 1999-148757P PRIORITY INFORMATION: 20000314 (60) US 2000-189026P

DOCUMENT TYPE:

Utility

APPLICATION

FILE SEGMENT: LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

22 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 11232

. component that may be treated, prevented, and/or diagnosed with LINE COUNT: the compositions of the invention include, but are not limited to, SUMM

type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia, Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin

dependent diabetes mellitis, and autoimmune inflammatory eye.

. . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, SUMM

and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus).

. neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated SUMM with (type I or type II) diabetes

mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases".

(glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.

(gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone;

alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as

L12 ANSWER 8 OF 115 USPATFULL on STN

PROGLYCEM.TM...

ACCESSION NUMBER:

2003:251152 USPATFULL

TITLE:

DETD

Serine protease polynucleotides, polypeptides, and

antibodies

INVENTOR (S):

Shi, Yanggu, Gaithersburg, MD, UNITED STATES

Ruben, Steven M., Brookeville, MD, UNITED STATES Ni, Jian, Germantown, MD, UNITED STATES

Young, Paul E., Gaithersburg, MD, UNITED STATES

Human Genome Sciences, Inc., Rockville, MD (U.S.

PATENT ASSIGNEE(S): corporation)

> KIND DATE NUMBER -----

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.: US 2003175938 A1 20030918 A1 20021216 (10) US 2002-319519

Continuation-in-part of Ser. No. US 2002-125459, filed on 19 Apr 2002, PENDING Continuation of Ser. No. US

2001-946633, filed on 6 Sep 2001, ABANDONED

Continuation of Ser. No. US 2000-597839, filed on 20 Jun 2000, ABANDONED Continuation-in-part of Ser. No. WO

2000-US12207, filed on 5 May 2000, PENDING Continuation-in-part of Ser. No. US 2000-597842, filed on 20 Jun 2000, ABANDONED Continuation-in-part of Ser. No. WO 2000-US12207, filed on 5 May 2000, PENDING

Continuation-in-part of Ser. No. US 2000-597843, filed on 20 Jun 2000, ABANDONED Continuation-in-part of Ser.

No. US 2002-67761, filed on 8 Feb 2002, PENDING Continuation of Ser. No. US 2001-804156, filed on 13 Mar 2001, ABANDONED

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DATE
                              NUMBER
                          _____
                       US 1999-133239P 19990507 (60)
PRIORITY INFORMATION:
                       US 1999-135163P 19990520 (60)
                       US 1999-147005P 19990803 (60)
                       US 1999-152935P 19990909 (60)
                       US 1999-162979P 19991101 (60)
                                          19990507 (60)
                       US 1999-133239P
                                          19990520 (60)
                       US 1999-135163P
                       US 1999-147005P
                                          19990803 (60)
                                           19990909 (60)
                       US 1999-152935P
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                       US 1999-162979P
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                       US 1999-147005P
                                           19990909 (60)
                       US 1999-152935P
                                           19991101 (60)
                        US 1999-162979P
                                           20000314 (60)
                        US 2000-189025P
                        US 2000-189025P 20000314 (60)
                        Utility
DOCUMENT TYPE:
                        APPLICATION
FILE SEGMENT:
                        HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
LEGAL REPRESENTATIVE:
                        ROCKVILLE, MD, 20850
                        22
NUMBER OF CLAIMS:
EXEMPLARY CLAIM:
LINE COUNT:
       . . . polypeptides corresponding to this gene and/or agonists and/or
                        13106
       antagonists thereof may be used to diagnose, prognose, treat, prevent,
SUMM
       and/or ameliorate type II diabetes
       mellitus (insulin resistant diabetes mellitus).
        . . . neutralizing or antagonistic antibodies) may be used to
       diagnose, prognose, treat, prevent, or ameliorate conditions associated
EUMM
       with (type I or type II) diabetes
       mellitus, including, but not limited to, diabetic
       ketoacidosis, diabetic coma, nonketotic hyperglycemic-
hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis,
       microvascular disease, hypertension, stroke, and other. . . renal
        failure, nephropathy other diseases and disorders as described in the
        "Renal Disorders" section), nerve damage, neuropathy, vision impairment
        (e.g., diabetic retinopathy and blindness), ulcers and
        impaired wound healing, infections (e.g., infectious diseases and
        disorders as described in the "Infectious Diseases". . .
           . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
 DETD
        (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
        AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
        such as rosiglitazone, AVANDIA.TM. (rosiglitazone
        maleate) ACTOS.TM. (piogliatazone), and troglitazone;
        alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
        such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
        PROGLYCEM.TM.. .
 L12 ANSWER 9 OF 115 USPATFULL on STN
                         2003:251148 USPATFULL
 ACCESSION NUMBER:
                         Protein tyrosine phosphatase polynucleotides,
 TITLE:
                         polypeptides, and antibodies
                         Shi, Yanggu, Gaithersburg, MD, UNITED STATES
 INVENTOR (S):
                         Ruben, Steven M., Olney, MD, UNITED STATES
                                                   DATE
                                           KIND
                              NUMBER
                                                 20030918
                                            A1
                         US 2003175934
 PATENT INFORMATION:
                                                20010824 (9)
                                           A1
                         US 2001-935703
 APPLICATION INFO :
                         Continuation-in-part of Ser. No. WO 2001-US5496, filed
 RELATED APPLN. INFO.:
                         on 22 Feb 2001, UNKNOWN
```

NUMBER

DATE

20000303 (60) US 2000-186658P PRIORITY INFORMATION: 20000316 (60) US 2000-189881P Utility DOCUMENT TYPE: APPLICATION HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, FILE SEGMENT: LEGAL REPRESENTATIVE: ROCKVILLE, MD, 20850 22 NUMBER OF CLAIMS: EXEMPLARY CLAIM: . component that may be treated, prevented, and/or diagnosed with LINE COUNT: the compositions of the invention include, but are not limited to, SUMM type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia, Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye. . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, SUMM and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus). . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated SUMM with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness,

cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.

(gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), DETD AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM..

L12 ANSWER 10 OF 115 USPATFULL on STN

2003:251072 USPATFULL ACCESSION NUMBER:

TITLE:

INVENTOR(S):

186 human secreted proteins

Ruben, Steven M., Olney, MD, UNITED STATES Rosen, Craig A., Laytonsville, MD, UNITED STATES Soppet, Daniel R., Centreville, VA, UNITED STATES Carter, Kenneth C., North Potomac, MD, UNITED STATES Bednarik, Daniel P., Columbia, MD, UNITED STATES Endress, Gregory A., Florence, MA, UNITED STATES Yu, Guo-Liang, Berkeley, CA, UNITED STATES Ni, Jian, Germantown, MD, UNITED STATES Feng, Ping, Germantown, MD, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Greene, John M., Gaithersburg, MD, UNITED STATES Ferrie, Ann M., Painted Post, NY, UNITED STATES Duan, D. Roxanne, Bethesda, MD, UNITED STATES Hu, Jing-Shan, Mountain View, CA, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES Olsen, Henrik S., Gaithersburg, MD, UNITED STATES Fischer, Carrie L., Burke, VA, UNITED STATES Ebner, Reinhard, Gaithersburg, MD, UNITED STATES Brewer, Laurie A., St. Paul, MN, UNITED STATES Moore, Paul A., Germantown, MD, UNITED STATES Shi, Yanggu, Gaithersburg, MD, UNITED STATES LaFleur, David W., Washington, DC, UNITED STATES

Li, Yi, Sunnyvale, CA, UNITED STATES Zeng, ZhiZhen, Lansdale, PA, UNITED STATES Kyaw, Hla, Frederick, MD, UNITED STATES

KIND

NUMBER

PATENT INFORMATION:	US 2003175858
APPLICATION INFO.:	US 2001-882171
RELATED APPLN. INFO.:	Continuation O
	Mar 2001. PEND

US 2003175858 A1 20030918
US 2001-882171 A1 20010618 (9)
Continuation of Ser. No. US 2001-809391, filed on 16
Mar 2001, PENDING Continuation of Ser. No. US
1998-149476, filed on 8 Sep 1998, GRANTED, Pat. No. US
6420526 Continuation-in-part of Ser. No. WO
1998-US4493, filed on 6 Mar 1998, PENDING

DATE

PRIORITY INFORMATION:

DATE NUMBER ______ 20000317 (60) US 2000-190068P 19970307 (60) US 1997-40162P 19970307 (60) US 1997-40333P 19970307 (60) US 1997-38621P 19970307 (60) US 1997-40626P 19970307 (60) US 1997-40334P 19970307 (60) US 1997-40336P 19970307 (60) US 1997-40163P 19970523 (60) US 1997-47600P 19970523 (60) US 1997-47615P 19970523 (60) US 1997-47597P 19970523 (60) US 1997-47502P 19970523 (60) US 1997-47633P 19970523 (60) US 1997-47583P 19970523 (60) US 1997-47617P 19970523 (60) US 1997-47618P 19970523 (60) US 1997-47503P 19970523 (60) US 1997-47592P 19970523 (60) US 1997-47581P 19970523 (60) US 1997-47584P 19970523 (60) US 1997-47500P 19970523 (60) US 1997-47587P 19970523 (60) US 1997-47492P 19970523 (60) US 1997-47598P 19970523 (60) US 1997-47613P 19970523 (60) US 1997-47582P 19970523 (60) US 1997-47596P 19970523 (60) US 1997-47612P 19970523 (60) US 1997-47632P US 1997-47601P 19970523 (60) 19970411 (60) US 1997-43580P 19970411 (60) US 1997-43568P 19970411 (60) US 1997-43314P 19970411 (60) US 1997-43569P 19970411 (60) US 1997-43311P 19970411 (60) US 1997-43671P 19970411 (60) US 1997-43674P 19970411 (60) US 1997-43669P 19970411 (60) US 1997-43312P US 1997-43313P 19970411 (60) 19970411 (60) US 1997-43672P 19970411 (60) US 1997-43315P 19970606 (60) US 1997-48974P 19970822 (60) US 1997-56886P 19970822 (60) US 1997-56877P 19970822 (60) US 1997-56889P US 1997-56893P 19970822 (60) 19970822 (60) US 1997-56630P 19970822 (60) US 1997-56878P 19970822 (60) US 1997-56662P 19970822 (60) US 1997-56872P 19970822 (60) US 1997-56882P 19970822 (60) US 1997-56637P

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                        US 1997-55724P
                        Utility
                        HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
                        APPLICATION
LEGAL REPRESENTATIVE:
                        ROCKVILLE, MD, 20850
                        23
                         2 Drawing Page(s)
                component that may be treated, prevented, and/or diagnosed with
       the compositions of the invention include, but are not limited to,
       type II collagen-induced arthritis, antiphospholipid
       syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
       polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
       polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
       pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
       dependent diabetes mellitus, and autoimmune inflammatory eye
                 polypeptides corresponding to this gene and/or agonists and/or
        antagonists thereof may be used to diagnose, prognose, treat, prevent,
        and/or ameliorate type II diabetes
        mellitus (insulin resistant diabetes mellitus).
             . neutralizing or antagonistic antibodies) may be used to
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DOCUMENT TYPE:

FILE SEGMENT:

LINE COUNT:

SUMM

SUMM

SUMM

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

disorders.

mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. DETD (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. . . [2309] The diabetic animals have many of the characteristic DETD features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al., J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol. 83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67 (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D. L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous diabetic mice develop hyperglycemia that is resistant to insulin analogous to human type II diabetes (Mandel et al., J Immunol. 120:1375-1377 (1978)).

diagnose, prognose, treat, prevent, and/or ameliorate conditions

associated with (type I or type II) diabetes

=> d 112 11-20 ibib, kwic

L12 ANSWER 11 OF 115 USPATFULL on STN

ACCESSION NUMBER:

2003:238753 USPATFULL

TITLE:

Trefoil domain-containing polynucleotides,

polypeptides, and antibodies

INVENTOR(S):

Ebner, Reinhard, Gaithersburg, MD, UNITED STATES Shi, Yanggu, Gaithersburg, MD, UNITED STATES Ruben, Steven M., Brookeville, MD, UNITED STATES

Human Genome Sciences, Inc., Rockville, MD, UNITED

STATES (U.S. corporation)

KIND DATE NUMBER -----

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.:

PATENT ASSIGNEE(S):

US 2003166911 A1 20030904 US 2002-266767 A1 20021009 (10) Continuation of Ser. No. US 2001-891171, filed on 26

Jun 2001, ABANDONED Continuation-in-part of Ser. No. WO

2000-US34920, filed on 22 Dec 2000, PENDING

DATE NUMBER ______

PRIORITY INFORMATION:

US 1999-171618P 19991223 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

22 1

LINE COUNT:

12173

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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . component that may be treated, prevented, and/or diagnosed with
       the compositions of the invention include, but are not limited to,
SUMM
       type II collagen-induced arthritis, antiphospholipid
       syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
       polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
       polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
       pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
       dependent diabetes mellitus, and autoimmune inflammatory eye
       disorders.
            . polypeptides corresponding to this gene and/or agonists and/or
       antagonists thereof may be used to diagnose, prognose, treat, prevent,
SUMM
       and/or ameliorate type II diabetes
       mellitus (insulin resistant diabetes mellitus).
            . neutralizing or antagonistic antibodies) may be used to
       diagnose, prognose, treat, prevent, or ameliorate conditions associated
SUMM
       with (type I or type II) diabetes
       mellitus, including, but not limited to, diabetic
       ketoacidosis, diabetic coma, nonketotic hyperglycemic-
       hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis,
       microvascular disease, hypertension, stroke, and other. . .
       failure, nephropathy other diseases and disorders as described in the
       "Renal Disorders" section), nerve damage, neuropathy, vision impairment
        (e.g., diabetic retinopathy and blindness), ulcers and
       impaired wound healing, infections (e.g., infectious diseases and
       disorders as described in the "Infectious Diseases".
              . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
DETD
        (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
       AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone
       maleate) ACTOS.TM. (piogliatazone), and troglitazone;
        alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
        such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
        PROGLYCEM.TM...
 L12 ANSWER 32 OF 115 USPATFULL on STN
                         2003:238383 USPATFULL
 ACCESSION NUMBER:
                         83 human secreted proteins
 TITLE:
                         Ruben, Steven M., Olney, MD, UNITED STATES
 INVENTOR (S):
                         Feng, Ping, Germantown, MD, UNITED STATES
                         LaFleur, David W., Washington, DC, UNITED STATES
                         Moore, Paul A., Germantown, MD, UNITED STATES
                         Shi, Yanggu, Gaithersburg, MD, UNITED STATES
                         Kyaw, Hla, Frederick, MD, UNITED STATES
                         Li, Yi, Sunnyvale, CA, UNITED STATES
                         Zeng, Zhizhen, Lansdale, PA, UNITED STATES
                         Carter, Kenneth C., North Potomac, MD, UNITED STATES
                         Endress, Gregory A., Florence, MA, UNITED STATES
                         Wei, Ying-Fei, Berkeley, CA, UNITED STATES
                         Fan, Ping, Potomac, MD, UNITED STATES
                         Rosen, Craig A., Laytonsville, MD, UNITED STATES
                         Human Genome Sciences, Inc., Rockville, MD, UNITED
 PATENT ASSIGNEE(S):
                         STATES, 20850 (U.S. corporation)
                                            KTND
                                                   DATE
                              MIMPED
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	MOMPHY		— ·	
PATENT INFORMATION:	US 2003166541		20030904	
APPLICATION INFO.: RELATED APPLN. INFO.:	US 2002-160162 Continuation-in-	part of	ED Contin	(10) US 1999-236557, filed uation-in-part of Ser. Jul 1998, PENDING
,	No. WO 1998-US15	949, Tl.	tea on 29	JUL 1990, PENDING

		NUMBER	DATE
PRIORITY	INFORMATION:	US 2001-295558P US 1997-54209P US 1997-54211P US 1997-54212P	20010605 (60) 19970730 (60) 19970730 (60) 19970730 (60)

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19970730 (60)
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                       US 1997-56730P
                       Utility
                       APPLICATION
                       HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
                       ROCKVILLE, MD, 20850
                        24
                        2 Drawing Page(s)
                        24088
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
         . . thrombocytopenia, idiopathic thrombocytopenia purpura, purpura
       (e.g., Henloch-Scoenlein purpura), autoimmunocytopenia, Goodpasture's
       syndrome, Pemphigus vulgaris, myasthenia gravis, Grave's disease
       (hyperthyroidism), and insulin-resistant diabetes mellitus.
       Additional disorders that are likely to have an autoimmune component
       that may be treated, prevented, and/or diagnosed with the compositions
       of the invention include, but are not limited to, type
       II collagen-induced arthritis, antiphospholipid syndrome,
       dermatitis, allergic encephalomyelitis, myocarditis, relapsing
       polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
       polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
       pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
       dependent diabetes mellitus, and autoimmune inflammatory eye
          . . component that may be treated, prevented, and/or diagnosed with
       the compositions of the invention include, but are not limited to,
       type II collagen-induced arthritis, antiphospholipid
       syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
       polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
       polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
       pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
       dependent diabetes mellitus, and autoimmune inflammatory eye
               polypeptides corresponding to this gene and/or agonists and/or
       antagonists thereof may be used to diagnose, prognose, treat, prevent,
       and/or ameliorate type II diabates
       mellitus (insulin resistant diabetes mellitus).
          . . neutralizing or antagonistic antibodies) may be used to
       diagnose, prognose, treat, prevent, and/or ameliorate conditions
        associated with (type I or type II) diabetes
        mellitus, including, but not limited to, diabetic
        ketoacidosis, diabetic coma, nonketotic hyperglycemic-
        hyperosmolar coma, seizures, mental confusion, drowsiness,
        cardiovascular disease (e.g., heart disease, atherosclerosis,
        microvascular disease, hypertension, stroke, and other. . .
        failure, nephropathy other diseases and disorders as described in the
        "Renal Disorders" section), nerve damage, neuropathy, vision impairment
        (e.g., diabetic retinopathy and blindness), ulcers and
        impaired wound healing, infections (e.g., infectious diseases and
        disorders as described in the "Infectious Diseases".
                 (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
        (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
        AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
```

DOCUMENT TYPE:

LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

disorders.

disorders.

FILE SEGMENT:

LINE COUNT:

DETD

DETD

DETD

DETD

DETD

```
such as rosiglitazone, AVANDIA.TM. (rosiglitazone
      maleate) ACTOS.TM. (piogliatazone), and troglitazone;
      alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
      such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
      [1679] The diabetic animals have many of the characteristic
      PROGLYCEM.TM..
      features observed in Type II diabetes
DETD
      mellitus. Homozygous (db+/db+) mice are obese in comparison to their
      normal heterozygous (db+/+m) littermates. Mutant diabetic
       (db+/db+) mice have a single autosomal recessive mutation on chromosome
       4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)).
       Animals show polyphagia, polydipsia and polyuria. Mutant
       diabetic mice (db+/db+) have elevated blood glucose, increased
       or normal insulin levels, and suppressed cell-mediated immunity (Mandel
       et al., J. Immunol.. . . glomerular filtration abnormalities have
       been described in these animals (Norido, F. et al., Exp. Neurol.
       83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67
       (1980); Giacomelli et Lab. Invest. 40(4):460-473 (1979); Coleman, D. L.,
       Diabetes 31 (Suppl):1 -6 (1982)). These homozygous
       diabetic mice develop hyperglycemia that is resistant to insulin
       analogous to human type II diabetes
       (Mandel et al., J. Immunol. 120:1375-1377 (1978)).
L12 ANSWER 13 OF 115 USPATFULL on STN
                        2003:237867 USPATFULL
                        Human G-protein chemokine receptor (CCR5) HDGNR10
ACCESSION NUMBER:
                        Rosen, Craig A., Laytonsville, MD, UNITED STATES
 TITLE:
                        Roschke, Viktor, Rockville, MD, UNITED STATES
 INVENTOR (S):
                        Li, Yi, Sunnyvale, CA, UNITED STATES
                        Ruben, Steven M., Olney, MD, UNITED STATES
                        Human Genome Sciences, Inc. (U.S. corporation)
 PATENT ASSIGNEE(S):
                                                  DATE
                            NUMBER KIND
                         ____________
                                           A1
                                                20030904
                         US 2003166024
                                         A1 20030501 (10)
A1 20020501 (10)
 PATENT INFORMATION:
                        US 2002-135839
 APPLICATION INFO.:
                         Continuation of Ser. No. US 2001-779879, filed on 9 Feb
 RELATED APPLN. INFO.:
                         2001, ABANDONED
                                             DATE
                               NUMBER
                         ______
                         US 2000-181258P 20000209 (60)
                         US 2000-234336P 20000922 (60)
Utility
 PRIORITY INFORMATION:
                         Utility
 DOCUMENT TYPE:
                         APPLICATION
                         STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C., 1100 NEW
 FILE SEGMENT:
 LEGAL REPRESENTATIVE:
                         YORK AVENUE, N.W., SUITE 600, WASHINGTON, DC,
                         20005-3934
                         61
 NUMBER OF CLAIMS:
  EXEMPLARY CLAIM:
                         4 Drawing Page(s)
  NUMBER OF DRAWINGS:
                         17941
  LINE COUNT:
  CAS INDEXING IS AVAILABLE FOR THIS PATENT.
           . . polypeptides corresponding to this gene and/or agonists and/or
         antagonists thereof may be used to diagnose, prognose, treat, prevent,
         and/or ameliorate type II diabetes
         mellitus (insulin resistant diabetes mellitus).
           . . neutralizing or antagonistic antibodies) may be used to
         diagnose, prognose, treat, prevent, and/or ameliorate conditions
  DETD
         associated with (type I or type II) diabetes
         mellitus, including, but not limited to, diabetic
         ketoacidosis, diabetic coma, nonketotic hyperglycemic-
         hyperosmolar coma, seizures, mental confusion, drowsiness,
         cardiovascular disease (e.g., heart disease, atherosclerosis,
         microvascular disease, hypertension, stroke, and other. . . described
         in the "Cardiovascular Disorders" section), dyslipidemia, kidney disease
          (e.g., renal failure and nephropathy) nerve damage, neuropathy, vision
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impairment (e.g., diabetic retinopathy and blindness), ulcers

```
and impaired wound healing, infections (e.g., infectious diseases and
      disorders as described in the "Infectious Diseases". . .
               (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRONTM
       (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
DETD
      AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone
      maleate) ACTOS.TM. (piogliatazone), and troglitazone;
       alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
       such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
       PROGLYCEM.TM.. .
       [1296] The diabetic animals have many of the characteristic
DETD
       features observed in Type II diabetes
       mellitus. Homozygous (db+/db+) mice are obese in comparison to their
       normal heterozygous (db+/+m) littermates. Mutant diabetic
       (db+/db+) mice have a single autosomal recessive mutation on chromosome
       4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)).
       Animals show polyphagia, polydipsia and polyuria. Mutant
       diabetic mice (db+/db+) have elevated blood glucose, increased
       or normal insulin levels, and suppressed cell-mediated immunity (Mandel
       et al, J. Immunol..... glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol.
       83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67
       (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.
       L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous
       diabetic mice develop hyperglycemia that is resistant to insulin
       analogous to human type II diabetes
       (Mandel et al, J. Immunol. 120:1375-1377 (1978)).
L12 ANSWER 14 OF 115 USPATFULL on STN
                        2003:232753 USPATFULL
ACCESSION NUMBER:
                        Attractin-like polynucleotides, polypeptides, and
TITLE:
                        antibodies
                        Ni, Jian, Germantown, MD, UNITED STATES
 INVENTOR (S):
                        Ruben, Steven M., Brookeville, MD, UNITED STATES
                        Young, Paul E., Gaithersburg, MD, UNITED STATES
                        Human Genome Sciences, Inc., Rockville, MD, UNITED
 PATENT ASSIGNEE(S):
                         STATES, 20850 (U.S. corporation)
                                          KIND DATE
                              NUMBER
                         -----
                         US 2003162954 A1 20030828 US 2002-193109 A1 20020712 (10)
 PATENT INFORMATION:
 APPLICATION INFO .:
                         Continuation of Ser. No. US 2002-84994, filed on 1 Mar
 RELATED APPLN. INFO.:
                         2002, ABANDONED Continuation of Ser. No. US
                         2001-790621, filed on 23 Feb 2001, ABANDONED
                         Continuation-in-part of Ser. No. WO 2000-US23663, filed
                         on 29 Aug 2000, PENDING
                               NUMBER DATE
                         ______
                         US 1999-151348P 19990830 (60)
 PRIORITY INFORMATION:
                         Utility
 DOCUMENT TYPE:
                         APPLICATION
 FILE SEGMENT:
                         HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
 LEGAL REPRESENTATIVE:
                         ROCKVILLE, MD, 20850
 NUMBER OF CLAIMS:
                         22
 EXEMPLARY CLAIM:
                         1
                         11989
 LINE COUNT:
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
        . . . component that may be treated, prevented, and/or diagnosed with
        the compositions of the invention include, but are not limited to,
 SUMM
        type II collagen-induced arthritis, antiphospholipid
        syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
        polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia,
        Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune
        Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin
        dependent diabetes mellitis, and autoimmune inflammatory eye.
         . . . polypeptides corresponding to this gene and/or agonists and/or
  SUMM
        antagonists thereof may be used to diagnose, prognose, treat, prevent,
```

```
and/or ameliorate type II diabetes
       mellitus (insulin resistant diabetes mellitus).
       . . . neutralizing or antagonistic antibodies) may be used to
       diagnose, prognose, treat, prevent, or ameliorate conditions associated
SUMM
       with (type I or type II) diabetes
       mellitus, including, but not limited to, diabetic
       ketoacidosis, diabetic coma, nonketotic hyperglycemic-
       hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal
       failure, nephropathy other diseases and disorders as described in the
       "Renal Disorders" section), nerve damage, neuropathy, vision impairment
       (e.g., diabetic retinopathy and blindness), ulcers and
       impaired wound healing, infections (e.g., infectious diseases and
       disorders as described in the "Infectious Diseases". .
          . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
        (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
DETD
       AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone;
        alpha-glucosidase inhibitors; bovine or porcine alucagon; somatostatins
        such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
        PROGLYCEM.TM..
 L12 ANSWER 15 OF 115 USPATFULL on STN
                         2003:231975 USPATFULL
                         NK cell receptor polynucleotides, polypeptides, and
 ACCESSION NUMBER:
 TITLE:
                          antibodies
                         Ruben, Steven M., Olney, MD, UNITED STATES
                          Shi, Yanggu, Gaithersburg, MD, UNITED STATES
 INVENTOR (S):
                                            KIND DATE
                               NUMBER
                          ______
                                             A1 20030328
                          US 2003162175 A1 20030328
US 2001-891464 A1 20010627 (9)
 PATENT INFORMATION:
                          Continuation-in-part of Ser. No. WO 2000-US34770, filed
 APPLICATION INFO .:
 RELATED APPLN. INFO .:
                          on 21 Dec 2000, UNKNOWN
                                 NUMBER
                                              DATE
                          -----
                          US 1999-171506P 19991222 (60)
  PRIORITY INFORMATION:
                          Utility
  DOCUMENT TYPE:
                          APPLICATION
                          HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
  FILE SEGMENT:
  LEGAL REPRESENTATIVE:
                          ROCKVILLE, MD, 20850
                          22
  NUMBER OF CLAIMS:
                           1
  EXEMPLARY CLAIM:
                           12365
  LINE COUNT:
  CAS INDEXING IS AVAILABLE FOR THIS PATENT.
         . . . component that may be treated, prevented, and/or diagnosed with
          the compositions of the invention include, but are not limited to,
  SUMM
          type II collagen-induced arthritis, antiphospholipid
          syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
          polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
          polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
          pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
          dependent diabetes mellitus, and autoimmune inflammatory eye
          . . . polypeptides corresponding to this gene and/or agonists and/or
          disorders.
          antagonists thereof may be used to diagnose, prognose, treat, prevent,
   SUMM
          and/or ameliorate type II diabetes
          mellitus (insulin resistant diabetes mellitus).
          . . neutralizing or antagonistic antibodies) may be used to
          diagnose, prognose, treat, prevent, or ameliorate conditions associated
   SUMM
          with (type I or type II) diabetes
          mellitus, including, but not limited to, diabetic
          ketoacidosis, diabetic coma, nonketotic hyperglycemic-
hyperosmolar coma, seizures, mental confusion, drowsiness,
          cardiovascular disease (e.g., heart disease, atherosclerosis,
```

microvascular disease, hypertension, stroke, and other. . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". . . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), DETD AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDS) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM. TM.. L12 ANSWER 16 OF 115 USPATFULL on STN 2003:219643 USPATFULL ACCESSION NUMBER: Human secreted proteins Barash, Steven C., Rockville, MD, UNITED STATES TITLE: INVENTOR (S): Ni, Jian, Germantown, MD, UNITED STATES Ruben, Steven M., Olney, MD, UNITED STATES Rosen, Craig A., Laytonsville, MD, UNITED STATES Shi, Yanggu, Gaithersburg, MD, UNITED STATES DATE NOMBER KIND DATE KIND US 2003152933 A1 20030814 US 2002-72977 A1 20020212 (10) PATENT INFORMATION: Continuation-in-part of Ser. No. WO 2001-US25288, filed APPLICATION INFO.: RELATED APPLN. INFO.: on 13 Aug 2001, PENDING DATE NUMBER -----US 2000-225215P 20000814 (60) PRIORITY INFORMATION: Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT: LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850 NUMBER OF CLAIMS: EXEMPLARY CLAIM: LINE COUNT: 12684 CAS INDEXING IS AVAILABLE FOR THIS PATENT. . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEMT.TM.. . . [0960] The diabetic animals have many of the characteristic DETD features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al., J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol. 83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67 (1980); Giacomelli et Lab Invest. 40(4):460-473 (1979); Coleman, D. L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous diabetic mice develop hyperglycemia that is resistant to insulin

analogous to human type II diabetes

(Mandel et al., J. Immunol. 120:1375-1377 (1978)).

ACCESSION NUMBER:

2003:214613 USPATFULL

TITLE:

Transferrin polynucleotides, polypeptides, and

antibodies

INVENTOR (S):

Ruben, Steven M., Brookeville, MD, UNITED STATES

Shi, Yanggu, Gaithersburg, MD, UNITED STATES

PATENT ASSIGNEE(S):

Human Genome Sciences, Inc., Rockville, MD, UNITED

STATES, 20850 (U.S. corporation)

NUMBER KIND -----

PATENT INFORMATION:

US 2003149256 A1 20030807 US 2002-266745 A1 20021009 (10)

RELATED APPLN. INFO.: Continuation of Ser. No. US 2001-891126, filed on 26 Jun 2001, PENDING Continuation-in-part of Ser. No. WO

2000-US34769, filed on 21 Dec 2000, PENDING

NUMBER DATE -----

PRIORITY INFORMATION: US 1999-171595P 19991223 (60)

DOCUMENT TYPE: Utility

APPLICATION

LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

LINE COUNT:

SUMM

DETD

12034

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . component that may be treated, prevented, and/or diagnosed with

the compositions of the invention include, but are not limited to,

type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin

dependent diabetes mellitus, and autoimmune inflammatory eye

disorders.

. . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent,

and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus).

. . neutralizing or antagonistic antibodies) may be used to SUMM

diagnose, prognose, treat, prevent, or ameliorate conditions associated

with (type I or type II) diabetes

mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemic-

hyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis,

microvascular disease, hypertension, stroke, and other. . . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment

(e.g., diabetic retinopathy and blindness), ulcers and

impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases".

. (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.

(gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)

such as rosiglitazone, AVANDIA.TM. (rosiglitazone

maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as

PROGLYCEM.TM.. .

L12 ANSWER 18 OF 115 USPATFULL on STN

2003:214304 USPATFULL ACCESSION NUMBER:

TITLE:

Major intrinsic protein (MIP)-like polynucleotides,

polypeptides and antibodies

Ruben, Steven M., Brookeville, MD, UNITED STATES INVENTOR(S):

Ni, Jian, Germantown, MD, UNITED STATES

Human Genome Sciences, Inc., Rockville, MD (U.S. PATENT ASSIGNEE(S):

corporation)

KIND DATE NUMBER ______ A1 20030807 US 2003148947 PATENT INFORMATION: Al 20020926 (10) US 2002-254939 Continuation of Ser. No. US 2001-862419, filed on 23 APPLICATION INFO.: RELATED APPLN. INFO.: May 2001, PENDING Continuation-in-part of Ser. No. WO 2000-US31919, filed on 21 Nov 2000, PENDING DATE NUMBER -----US 1999-167247P 19991124 (60) PRIORITY INFORMATION: Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, LEGAL REPRESENTATIVE: ROCKVILLE, MD, 20850 NUMBER OF CLAIMS: EXEMPLARY CLAIM: . 11742 LINE COUNT: CAS INDEXING IS AVAILABLE FOR THIS PATENT. . . . component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye . . . polypeptides corresponding to this gene and/or agonists and/or disorders. antagonists thereof may be used to diagnose, prognose, treat, prevent, SUMM and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus). . . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated SUMM with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". . . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), DETD AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. . L12 ANSWER 19 OF 115 USPATFULL on STN 2003:187895 USPATFULL ACCESSION NUMBER: 12 human secreted proteins Ni, Jian, Germantown, MD, UNITED STATES TITLE: Young, Paul E., Gaithersburg, MD, UNITED STATES INVENTOR (S): Kenny, Joseph J., Damascus, MD, UNITED STATES Olsen, Henrik S., Gaithersburg, MD, UNITED STATES Moore, Paul A., Germantown, MD, UNITED STATES

KIND DATE

Wei, Ying-Fei, Berkeley, CA, UNITED STATES Greene, John M., Gaithersburg, MD, UNITED STATES Ruben, Steven M., Olney, MD, UNITED STATES

NUMBER

A1 20030710 US 2003129685 PATENT INFORMATION: A1 20010418 (9) US 2001-836353 Continuation-in-part of Ser. No. WO 1999-US25031, filed APPLICATION INFO.: RELATED APPLN. INFO.: on 27 Oct 1999, UNKNOWN DATE NUMBER -----19981028 (60) US 1998-105971P PRIORITY INFORMATION: US 2000-198407P 20000419 (60) Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, LEGAL REPRESENTATIVE: ROCKVILLE, MD, 20850 23 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 59 Drawing Page(s) NUMBER OF DRAWINGS: 31945. LINE COUNT: CAS INDEXING IS AVAILABLE FOR THIS PATENT. . . component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye . . polypeptides corresponding to this gene and/or agonists and/or disorders. antagonists thereof may be used to diagnose, prognose, treat, prevent, DETD and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus). . . . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, and/or ameliorate conditions DETD associated with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), DETD AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. . [1496] The diabetic animals have many of the characteristic DETD features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al., J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol.

.83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67

diabetic mice develop hyperglycemia that is resistant to insulin

L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous

analogous to human type 11 diabetes (Mandel et al., J.

(1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.

L12 ANSWER 20 OF 115 USPATFULL on STN 2003:173232 USPATFULL ACCESSION NUMBER: TITLE:

INVENTOR (S):

B7-like polynucleotides, polypeptides, and antibodies

Ruben, Steven M., Olney, MD, UNITED STATES Chen, Lieping, Rochester, MN, UNITED STATES Baker, Kevin P., Darnestown, MD, UNITED STATES

Ni, Jian, Germantown, MD, UNITED STATES

Human Genome Sciences, Inc., Rockville, MD, UNITED PATENT ASSIGNEE(S):

STATES, 20850 (U.S. corporation)

KIND NUMBER -----US 2003119076 A1 20030626 US 2002-141953 A1 20020510 (10) PATENT INFORMATION:

APPLICATION INFO.: Continuation of Ser. No. US 2001-790622, filed on 23

Feb 2001, PENDING Continuation-in-part of Ser. No. WO RELATED APPLN. INFO.:

2000-US23792, filed on 30 Aug 2000, PENDING

DATE NUMBER

US 1999-152317P 19990903 (60) PRIORITY INFORMATION: US 2000-200346P 20000428 (60)

Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, LEGAL REPRESENTATIVE:

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

15 Drawing Page(s) NUMBER OF DRAWINGS:

12418 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to. DETD type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia, Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye.

. . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, DETD and/or ameliorate type II diabetes

mellitus (insulin resistant diabetes mellitus).

. . . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated DETD with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and

disorders as described in the "Infectious Diseases". . . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)

such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. .

DETD

L12 ANSWER 21 OF 115 USPATFULL on STN 2003:165984 USPATFULL ACCESSION NUMBER: 25 human secreted proteins TITLE: Rosen, Craig A., Laytonsville, MD, UNITED STATES INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES Fiscella, Michele, Bethesda, MD, UNITED STATES Wei, Ping, Brookeville, MD, UNITED STATES Baker, Kevin P., Darnestown, MD, UNITED STATES Birse, Charles E., North Potomac, MD, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Komatsoulis, George A., Silver Spring, MD, UNITED STATES Moore, Paul A., Germantown, MD, UNITED STATES Soppet, Daniel R., Centreville, VA, UNITED STATES Human Genome Sciences, Inc., Rockville, MD, UNITED PATENT ASSIGNEE(S): STATES, 20850 (U.S. corporation) DATE KIND NUMBER -----US 2003113840 A1 20030619 PATENT INFORMATION: Al 20020201 (10) US 2002-60255 APPLICATION INFO.: Continuation of Ser. No. US 2001-781417, filed on 13 RELATED APPLN. INFO.: Feb 2001, ABANDONED Continuation-in-part of Ser. No. WO 2000-US22325, filed on 16 Aug 2000, PENDING DATE NUMBER US 1999-149182P 19990817 (60) PRIORITY INFORMATION: Utility DOCUMENT TYPE: APPLICATION FILE SEGMENT: HUMAN GENOME SCIENCES INC. 9410 KEY WEST AVENUE, LEGAL REFRESENTATIVE: ROCKVILLE, MD, 20850 NUMBER OF CLAIMS: 1 EXEMPLARY CLAIM: 20339 LINE COUNT: CAS INDEXING IS AVAILABLE FOR THIS PATENT. SUMM . . . autosomal recessive, childhood-onset Retinitis pigmentosa-20 Hypothyroidism, nongoitrous 188540 [Urate oxidase deficiency] 191540 Acyl-CoA dehydrogenase, medium chain, deficiency of 201450 Maple syrup urine disease, type II 248610 Galactosialidosis 256540 Thymine-uraciluria 274270 Fluorouracil toxicity, sensitivity to HMG-CoA synthease-2 deficiency 500234 MODY, type 1, 125850 600281 Non-insulin-dependent diabetes mellitus, 125853 Atrioventricular canal defect-1 600309 Retinitis pigmentosa-18 501414 Acute insulin response 601676 Cone-rod dystrophy 3 501691 Fundus flavimaculatus with macular dystrophy, 248200 Retinitis. . . . component that may be treated, prevented, and/or diagnosed with SUMM the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders. . and/or polypeptides corresponding to this gene and/or agonists SUMM and/or antagonists thereof maybe used to diagnose, prognose, treat, prevent, and/or ameliorate type II diabetes

mellitus (insulin resistant diabetes mellitus).

```
. . neutralizing or antagonistic antibodies) may be used to
SUMM .
      diagnose, prognose, treat, prevent, and/or ameliorate conditions
      associated with (type I or type II) diabetes
      mellitus, including, but not limited to, diabetic
      ketoacidosis, diabetic coma, nonketotic hyperglycemic-
      hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis,
       microvascular disease, hypertension, stroke, and other. . . renal
       failure, nephropathy other diseases and disorders as described in the
       "Renal Disorders" section), nerve damage, neuropathy, vision impairment
       (e.g., diabetic retinopathy and blindness), ulcers and
       impaired wound healing, infections (e.g., infectious diseases and
       disorders as described in the "Infectious Diseases". .
         . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
DETD
       (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
       AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone
       maleate) ACTOS.TM. (piogliatazone), and troglitazone;
       alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
       such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
       PROGLYCEM.TM.. .
       [1215] The diabetic animals have many of the characteristic
DETD
       features observed in Type II diabetes
       mellitus. Homozygous (db+/db+) mice are obese in comparison to their
       normal heterozygous (db+/+m) littermates. Mutant diabetic
       (db+/db+) mice have a single autosomal recessive mutation on chromosome
       4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)).
       Animals show polyphagia, polydipsia and polyuria. Mutant
       diabetic mice (db+/db+) have elevated blood glucose, increased
       or normal insulin levels, and suppressed cell-mediated immunity (Mandel
       et al., J. Immunol.. . . glomerular filtration abnormalities have
       been described in these animals (Norido, F. et al., Exp. Neurol.
       83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67
       (1930); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.
       L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous
       diabetic mice develop hyperglycemia that is resistant to insulin
        analogous to human type II diabetes
        (Mandel et al., J. Immunol. 120:1375-1377 (1978)).
L12 ANSWER 22 OF 115 USPATFULL on STN
                         2003:146312 USPATFULL
ACCESSION NUMBER:
                       Human G-protein Chemokine Receptor (CCR5) HDGNR10
TITLE:
                         Roschke, Viktor, Rockville, MD, UNITED STATES
 INVENTOR(S):
                         Rosen, Craig A., Laytonsville, MD, UNITED STATES
                         Ruben, Steven M., Olney, MD, UNITED STATES
Human Genome Sciences, Inc. (U.S. corporation)
 PATENT ASSIGNEE(S):
                                       KIND DATE
                            NUMBER
                      . -----
 PATENT INFORMATION: US 2003100058 A1 20030529
APPLICATION INFO:: US 2002-67800 A1 20020208 (10)
                         Continuation-in-part of Ser. No. WO 2001-US4153, filed
 RELATED APPLN. INFO.:
                         on 9 Feb 2001, UNKNOWN Continuation-in-part of Ser. No.
                         US 2001-779880, filed on 9 Feb 2001, PENDING
                                             DATE
                                NUMBER
                         -----
                         US 2001-297257P 20010612 (60)
 PRIORITY INFORMATION:
                         US 2001-310458P 20010808 (60)
                         US 2001-328447P 20011012 (60)
US 2001-341725P 20011221 (60)
 DOCUMENT TYPE:
                         Utility
                         APPLICATION
 FILE SEGMENT:
                         STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C., 1100 NEW
 LEGAL REPRESENTATIVE:
                         YORK AVENUE, N.W., SUITE 600, WASHINGTON, DC,
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20005-3934

7 Drawing Page(s)

60

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

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18955
LINE COUNT:
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . polypeptides corresponding to this gene and/or agonists and/or
      antagonists thereof may be used to diagnose, prognose, treat, prevent,
       and/or ameliorate type II diabetes
       mellitus (insulin resistant diabetes mellitus).
         . . neutralizing or antagonistic antibodies) may be used to
DETD
       diagnose, prognose, treat, prevent, and/or ameliorate conditions
      associated with (type I or type II) diabetes
       mellitus, including, but not limited to, diabetic
       ketoacidosis, diabetic coma, nonketotic hyperglycemic-
       hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis,
       microvascular disease, hypertension, stroke, and other. . . described
       in the "Cardiovascular Disorders" section), dyslipidemia, kidney disease
       (e.g., renal failure and nephropathy) nerve damage, neuropathy, vision
       impairment (e.g., diabetic retinopathy and blindness), ulcers
       and impaired wound healing, infections (e.g., infectious diseases and
       disorders as described in the "Infectious Diseases".
        . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
DETD
       (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
       AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone
       maleate) ACTOS.TM. (piogliatazone), troglitazone;
       alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
       such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
       PROGLYCEM.TM. (diazoxide) .. .
       [1355] The diabetic animals have many of the characteristic
DETD
       features observed in Type II diabetes
       mellitus. Homozygous (db+/db+) mice are obese in comparison to their
       normal heterozygous (db+/+m) littermates. Mutant diabetic
       (db+/db+) mice have a single autosomal recessive mutation on chromosome
       4 (db+) (Coleman et al. Proc. Natl Acad. Sci. USA 77:283-293 (1982)).
       Animals show polyphagia, polydipsia and polyuria. Mutant
       diabetic mice (db+/db+) have elevated blood glucose, increased
       or normal insulin levels, and suppressed cell-mediated immunity (Mandel
       et al., J. Immunol.. . . glomerular filtration abnormalities have
       been described in these animals (Norido, F. et al., Exp. Neurol.
       83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67
       (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.
       L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous
       diabetic mice develop hyperglycemia that is resistant to insulin
       analogous to human type II diabetes
        (Mandel et al., J. Immunol. 120:1375-1377 (1978)).
L12 ANSWER 23 OF 115 USPATFULL on STN
                        2003:120997 USPATFULL
ACCESSION NUMBER:
                        25 human prostate and prostate cancer associated
TITLE:
                        proteins
                        Birse, Charles E., North Potomac, MD, UNITED STATES
INVENTOR (S):
                        Rosen, Craig A., Laytonsville, MD, UNITED STATES
                        Human Genome Sciences, Inc., Rockville, MD, UNITED
PATENT ASSIGNEE(S):
                        STATES, 20850 (U.S. corporation)
                                       KIND DATE
                             NUMBER
                        _____
                        US 2003083481 A1 20030501
US 2002-36542 A1 20020107 (10)
PATENT INFORMATION:
APPLICATION INFO.:
                        Continuation-in-part of Ser. No. WO 2000-US19666, filed
RELATED APPLN. INFO.:
                        on 20 Jul 2000, UNKNOWN
                                            DATE
                               NUMBER
                        US 1999-144972P 19990721 (60)
PRIORITY INFORMATION:
                        US 1999-148681P 19990813 (60)
                        US 1999-149173P 19990817 (60)
                        US 1999-158004P 19991006 (60)
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US 2000-194689P 20000405 (60)

Utility

DOCUMENT TYPE:

```
APPLICATION
FILE SEGMENT:
                       HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
LEGAL REPRESENTATIVE:
                       ROCKVILLE, MD, 20850
NUMBER OF CLAIMS:
EXEMPLARY CLAIM:
                       26241
LINE COUNT:
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
         . . tissues and organs (e.g., cancers of the hypothalamus,
       pituitary gland, thyroid gland, parathyroid glands, pancreas, adrenal
       glands, ovaries, and testes), diabetes (e.g., diabetes
       insipidus, type I and type II diabetes
       mellitus), obesity, disorders related to pituitary glands (e.g.,
       hyperpituitarism, hypopituitarism, and pituitary dwarfism),
       hypothyroidism, hyperthyroidism, goiter, reproductive disorders (e.g.
                . adrenal glands (e.g., Addison's Disease, corticosteroid
       deficiency, and Cushing's Syndrome), kidney cancer (e.g., hypemephroma,
       transitional cell cancer, and Wilm's tumor), diabetic
       nephropathy, interstitial nephritis, polycystic kidney disease,
       glomerulonephritis (e.g., IgM mesangial proliferative glomerulonephritis
       and glomerulonephritis caused by autoimmune disorders; such as. .
                component that may be treated, prevented, and/or diagnosed with
SUMM
       the compositions of the invention include, but are not limited to,
       type II collagen-induced arthritis, antiphospholipid
       syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
       polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
       polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
       pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
       dependent diabetes mellitus, and autoimmune inflammatory eye
       disorders.
      . . . polypeptides corresponding to this gene and/or agonists and/or
SUMM
       antagonists thereof may be used to diagnose, prognose, treat, prevent,
       and/or ameliorate type II diabetes
       mellitus (insulin resistant diabetes mellitus).
       . . neutralizing or antagonistic antibodies) may be used to
SUMM
       diagnose, prognose, treat, prevent, and/or ameliorate conditions
       associated with (type I or type II) diabetes
       mellitus, including, but not limited to, diabetic
       ketoacidosis, diabetic coma, nonketotic hyperglycemic-
       hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis,
       microvascular disease, hypertension, stroke, and other. . renal
       failure, nephropathy other diseases and disorders as described in the
       "Renal Disorders" section), nerve damage, neuropathy, vision impairment
       (e.g., diabetic retinopathy and blindness), ulcers and
       impaired wound healing, infections (e.g., infectious diseases and
       disorders as described in the "Infectious Diseases".
                (glyburide), GLUCOTRIL.TM. (glipizide), and DIAMICRON.TM.
DETD
       (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
       AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone
       maleate) ACTOS.TM. (piogliatazone), and troglitazone;
       alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
       such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
       PROGLYCEM.TM..
        [1380] The diabetic animals have many of the characteristic
DETD
       features observed in Type II diabetes
       mellitus. Homozygous (db+/db+) mice are obese in comparison to their
       normal heterozygous (db+/+m) littermates. Mutant diabetic
        (db+/db+) mice have a single autosomal recessive mutation on chromosome
       4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)).
       Animals show polyphagia, polydipsia and polyuria. Mutant
       diabetic mice (db+/db+) have elevated blood glucose, increased
       or normal insulin levels, and suppressed cell-mediated immunity (Mandel
       et al., J. Immunol.. . . glomerular filtration abnormalities have
       been described in these animals (Norido, F. et al., Exp. Neurol.
       83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67
        (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.
       L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous
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diabetic mice develop hyperglycemia that is resistant to insulin

analogous to human type II diabetes (Mandel et al., J. Immunol. 120:1375-1377 (1978)).

L12 ANSWER 24 OF 115 USPATFULL on STN

ACCESSION NUMBER:

INVENTOR(S):

2003:79163 USPATFULL

TITLE:

Combination therapy comprising glucose reabsorption

inhibitors and retinoid-X receptor modulators

Bussolari, Jacqueline C., Skillman, NJ, UNITED STATES

Chen, Xiaoli, Belle Mead, NJ, UNITED STATES Conway, Bruce R., Doylestown, PA, UNITED STATES Demarest, Keith T., Flemington, NJ, UNITED STATES Ross, Hamish N.M., Far Hills, NJ, UNITED STATES

Severino, Rafael, Madrid, SPAIN

KIND NUMBER

PATENT INFORMATION: APPLICATION INFO.:

US 2003055091 A1 US 2002-115725 A1 20030320

20020403 (10)

NUMBER DATE

PRIORITY INFORMATION:

US 2001-281479P 20010404 (60)

DOCUMENT TYPE: FILE SEGMENT:

Utility

APPLICATION

LEGAL REPRESENTATIVE:

AUDLEY A. CIAMPORCERO JR., JOHNSON & JOHNSON, ONE

JOHNSON & JOHNSON PLAZA, NEW BRUNSWICK, NJ, 08933-7003

NUMBER OF CLAIMS:

1

EXEMPLARY CLAIM: LINE COUNT:

2308

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

[0005] Type II diabetes mellitus

(non-insulin-dependent diabetes mellitus or NIDDM) is a metabolic disorder involving the dysregulation of glucose metabolism and

impaired insulin sensitivity. Type II

diabetes mellitus usually develops in adulthood and is associated with the body's inability to utilize or make sufficient insulin. In addition to the insulin resistance observed in the target

tissues, patients suffering from type II

diabetes mellitus have a relative insulin deficiency--that is, patients have lower than predicted insulin levels for a given plasma

glucose concentration. Type II diabetes

mellitus is characterized by the following clinical signs or symptoms: persistently elevated plasma glucose concentration or hyperglycemia;

polyuria; polydipsia and/or.

. . recognized in some 2% of diagnostic coronary catheterizations. SUMM

Often disabling, it presents symptoms or risk factors for the

development of Type II diabetes mellitus

and cardiovascular disease, including impaired glucose tolerance (IGT), impaired fasting glucose (IFG), hyperinsulinemia, insulin resistance,

dyslipidemia (e.g., high triglycerides,. .

[0008] Typical treatment of Type II diabetes SUMM

mellitus focuses on maintaining the blood glucose level as near to normal as possible with lifestyle modification relating to diet and exercise, and when necessary, the treatment with anti-diabetic agents, insulin or a combination thereof. NIDDM that cannot be controlled by dietary management is treated with oral antidiabetic

agents.

SUMM

therapies typically include metformin and sulfonylureas as well as thiazolidinediones. Metformin monotherapy is a first line choice,

particularly for treating type II diabetic

patients who are also obese and/or dyslipidemic. Lack of an appropriate response to metformin is often followed by treatment with. . . Alpha glucosidase inhibitors are also used as first and second line therapies. Patients who do not respond appropriately to oral anti-diabetic

monotherapy, are given combinations of the above-mentioned agents. When glycemic control cannot be maintained with oral antidiabetics alone, insulin therapy.

[0019] U.S. Pat. No. 6,153,632 to R. Rieveley discloses a method and DRWD composition stated to be for the treatment of diabetes

mellitus (Type I, Impaired Glucose Tolerance ["IGT"] and Type II), which incorporates a therapeutic amount of one or more insulin sensitizers along with one or more of an orally ingested insulin, an injected insulin, a sulfonylurea, a biguanide or an alpha-glucosidase inhibitor for the treatment of diabetes mellitus.

DRWD

. or 5-((4-(2-(methyl-2-pyridinylamino) ethoxy) phenyl) methyl)-2,4-thiazolidinedione, known as AVANDIA; also known as BRL 49653, BRL 49653C, BRL 49653C, SB 210232, or rosiglitazone maleate);

[0395] Thus, for treating diabetes, particularly Type DETD II diabetes mellitus, or Syndrome X, a compound of Formulae I, II, III, IV, or V in combination with one or more. formula I in the range of about 25 to 1000 mg once or twice daily and repeated doses of the anti-diabetic agent or agents at jointly effective dosages. The jointly effective dosage for RXR modulators disclosed herein may be readily determined.

L12 ANSWER 25 OF 115 USPATFULL on STN

ACCESSION NUMBER:

2003:72168 USPATFULL 64 human secreted proteins

TITLE: INVENTOR (S):

Ruben, Steven M., Olney, MD, UNITED STATES Rosen, Craig A., Laytonsville, MD, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Greene, John M., Gaithersburg, MD, UNITED STATES Ni, Jian, Germantown, MD, UNITED STATES

Feng, Ping, Gaithersburg, MD, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES Hu, Jing-Shan, Mountain View, CA, UNITED STATES Ferrie, Ann M., Tewksbury, MA, UNITED STATES Yu, Guo-Liang, Berkeley, CA, UNITED STATES

KIND DATE

Duan, Roxanne D., Bethesda, MD, UNITED STATES Janat, Fouad, Westerly, RI, UNITED STATES

------PATENT INFORMATION: APPLICATION INFO.:

RELATED APPLN. INFO.:

US 2003050455 20030313 A1 20010206 (9) US 2001-776724 A1

Continuation-in-part of Ser. No. US 2000-669688, filed

on 26 Sep 2000, PENDING Continuation of Ser. No. US

1999-229982, filed on 14 Jan 1999, PENDING Continuation-in-part of Ser. No. WO 1998-US14613, filed

on 15 Jul 1998, UNKNOWN

NUMBER

DATE NUMBER 20000208 (60) US 2000-180909P PRIORITY INFORMATION: 19970722 (60) US 1997-53442P 19970818 (60) US 1997-56359P 19970716 (60) US 1997-52661P 19970716 (60) US 1997-52872P 19970716 (60) US 1997-52871P 19970716 (60) US 1997-52874P 19970716 (60) US 1997-52873P 19970716 (50) US 1997-52870P 19970716 (60) US 1997-52875P 19970722 (60) US 1997-53440P 19970722 (60) US 1997-53441P Utility

DOCUMENT TYPE:

APPLICATION FILE SEGMENT:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, LEGAL REPRESENTATIVE:

ROCKVILLE, MD, 20850

23 NUMBER OF CLAIMS:

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

2 Drawing Page(s)

21934 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to,

type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders.

. . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, and/or ameliorate type II diabetes mellitus).

. . . neutralizing or antagonistic antibodies) may be used to

diagnose, prognose, treat, prevent, and/or ameliorate conditions associated with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemic-hyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases".

(glipizide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagen; somanostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM...

[1629] The diabetic animals have many of the characteristic features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+ m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al, J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol. 83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67 (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous diabetic mice develop hyperglycemia that is resistant to insulin analogous to human type II diabetes (Mandel et al., J Immunol. 120:1375-1377 (1978)).

L12 ANSWER 26 OF 115 USPATFULL on STN

ACCESSION NUMBER:

TITLE: INVENTOR(S):

DETD

DETD

DETU

DETD

2003:71333 USPATFULL 186 human secreted proteins Ruben, Steven M., Olney, MD, UNITED STATES Rosen, Craig A., Laytonsville, MD, UNITED STATES Soppet, Daniel R., Centreville, VA, UNITED STATES Carter, Kenneth C., North Potomac, MD, UNITED STATES Bednarik, Daniel P., Columbia, MD, UNITED STATES Endress, Gregory A., Florence, MA, UNITED STATES Yu, Guo-Liang, Berkeley, CA, UNITED STATES Ni, Jian, Germantown, MD, UNITED STATES Feng, Ping, Gaithersburg, MD, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Greene, John M., Gaithersburg, MD, UNITED STATES Ferrie, Ann M., Painted Post, NY, UNITED STATES Duan, D. Roxanne, Bethesda, MD, UNITED STATES Hu, Jing-Shan, Mountain View, CA, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES Olsen, Henrik S., Gaithersburg, MD, UNITED STATES

Fischer, Carrie L., Burke, VA, UNITED STATES Ebner, Reinhard, Gaithersburg, MD, UNITED STATES Brewer, Laurie A., St. Paul, MN, UNITED STATES Moore, Paul A., Germantown, MD, UNITED STATES Shi, Yanggu, Gaithersburg, MD, UNITED STATES LaFleur, David W., Washington, DC, UNITED STATES Li, Yi, Sunnyvale, CA, UNITED STATES Zeng, Zhizhen, Lansdale, PA, UNITED STATES Kyaw, Hla, Frederick, MD, UNITED STATES

KIND

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.:

NUMBER A1 20030313 US 2003049618 US 2001-809391 A1 20010316 (9) Continuation-in-part of Ser. No. US 1998-149476, filed on 8 Sep 1998, GRANTED, Pat. No. US 6420526 Continuation-in-part of Ser. No. WO 1998-US4493, filed on 6 Mar 1998, UNKNOWN

DATE

NUMBER

PRIORITY INFORMATION:

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US 1997-55724P
 Utility
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DOCUMENT TYPE:

FILE SEGMENT:

LEGAL REPRESENTATIVE:

APPLICATION

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 23 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS:

2 Drawing Page(s)

26235 LINE COUNT: DETD

component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders.

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antagonists thereof may be used to diagnose, prognose, treat, prevent,
       and/or ameliorate type II diabetes
       mellitus (insulin resistant diabetes mellitus).
          . . neutralizing or antagonistic antibodies) may be used to
DETD
       diagnose, prognose, treat, prevent, and/or ameliorate conditions
       associated with (type I or type II) diabetes
       mellitus, including, but not limited to, diabetic
       ketoacidosis, diabetic coma, nonketotic hyperglycemic-
       hyperosmolar coma, seizures, mental confusion, drowsiness,
       cardiovascular disease (e.g., heart disease, atherosclerosis,
       microvascular disease, hypertension, stroke, and other. . . renal
       failure, nephropathy other diseases and disorders as described in the
       "Renal Disorders" section), nerve damage, neuropathy, vision impairment
       (e.g., diabetic retinopathy and blindness), ulcers and
       impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases".
         . . (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
DETD
       (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
       AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
       such as rosiglitazone, AVANDIA.TM. (rosiglitazone
       maleate) ACTOS.TM. (piogliatazone), and troglitazone;
       alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
       such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
       PROGLYCEM.TM....
       [2107] The diabetic animals have many of the characteristic
DETD
       features observed in Type II diabetes
       mellitus. Homozygous (db+/db+) mice are obese in comparison to their
       normal heterozygous (db+/+m) littermates. Mutant diabetic
       (db+/db+) mice have a single autosomal recessive mutation on chromosome
       4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)).
       Animals show polyphagia, polydipsia and polyuria. Mutant
       diabetic mice (db+/db+) have elevated blood glucose, increased
       or normal insulin levels, and suppressed cell-mediated immunity (Mandel
       et al., J. Immunol.. . . glomerular filtration abnormalities have
       been described in these animals (Norido, F. et al., Exp. Neurol.
       83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67
       (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D.
       L., Diabetes 31 (Suppl):1-6 (1982)). These homozygous
       diabetic mice develop hyperglycemia that is resistant to insulin
       analogous to human type II diabetes
       (Mandel et al., J. Immunol. 120:1375-1377 (1978)).
L12 ANSWER 27 OF 115 USPATFULL on STN
                        2003:65429 USPATFULL
ACCESSION NUMBER:
                        Combination therapy comprising glucose reabsorption
TITLE:
                        inhibitors and PPAR modulators
                        Bussolari, Jacqueline C., Skillman, NJ, UNITED STATES Chen, Xiaoli, Belle Mead, NJ, UNITED STATES
INVENTOR(S):
                        Conway, Bruce R., Doylestown, PA, UNITED STATES
                        Demarest, Keith T., Flemington, NJ, UNITED STATES
                        Ross, Hamish N.M., Far Hills, NJ, UNITED STATES
                         Severino, Rafael, Madrid, SPAIN
                                         KIND
                             NUMBER
                                                   DATE
                        US 2003045553 A1
                                                 20030306
PATENT INFORMATION:
APPLICATION INFO.:
                        US 2002-115827
                                           A1
                                                 20020403 (10)
                                            DATE
                               NUMBER
                        US 2001-281429P
                                            20010404 (60)
PRIORITY INFORMATION:
DOCUMENT TYPE:
                        Utility
                        APPLICATION
FILE SEGMENT:
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AUDLEY A. CIAMPORCERO JR., JOHNSON & JOHNSON, ONE

JOHNSON & JOHNSON PLAZA, NEW BRUNSNICK, NJ, 08933-7003

. . . polypeptides corresponding to this gene and/or agonists and/or

DETD

LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

67

7 Drawing Page(s)

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2106
LINE COUNT:
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       [0005] Type II diabetes mellitus
SUMM
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(non-insulin-dependent diabetes mellitus or NIDDM) is a metabolic disorder involving the dysregulation of glucose metabolism and impaired insulin sensitivity. Type II diabetes mellitus usually develops in adulthood and is associated with the body's inability to utilize or make sufficient insulin. In addition to the insulin resistance observed in the target tissues, patients suffering from type II diabetes mellitus have a relative insulin deficiency--that is, patients have lower than predicted insulin levels for a given plasma glucose concentration. Type II diabetes

mellitus is characterized by the following clinical signs or symptoms: persistently elevated plasma glucose concentration or hyperglycemia; polyuria; polydipsia and/or. recognized in some 2% of diagnostic coronary catheterizations.

Often disabling, it presents symptoms or risk factors for the development of Type II diabetes mellitus and cardiovascular disease, including impaired glucose tolerance (IGT), impaired fasting glucose (IFG), hyperinsulinemia, insulin resistance, dyslipidemia (e.g., high triglycerides,.

[0008] Typical treatment of Type II diabetes mellitus focuses on maintaining the blood glucose level as near to SUMM normal as possible with lifestyle modification relating to diet.

therapies typically include metformin and sulfonylureas as well as thiazolidinediones. Metformin monotherapy is a first line choice, SUMM particularly for treating type II diabetic patients who are also obese and/or dyslipidemic. Lack of an appropriate response to metformin is often followed by treatment with.

[0027] US Pat. No. 6,153,632 to R. Rieveley discloses a method and composition stated to be for the treatment of diabetes mellitus (Type I, Impaired Glucose Tolerance ["IGT"] and Type II), which incorporates a therapeutic amount of one or more insulin sensitizers along with one or more of an orally ingested insulin, an injected insulin, a sulfonylurea, a biguanide or an alpha-glucosidase inhibitor for the treatment of diabetes

mellitus. or 5-((4-(2-(methyl-2-pyridinylamino) ethoxy) phenyl) methyl) -2,4-thiazolidinedione, known as AVANDIA; also known as BRL 49653, BRL 49653C, BRL 49653C, SB 210232, or rosiglitazone DETD maleate);

[0350] Thus, for treating diabetes, particularly Type DETD II diabetes mellitus, or Syndrome X, a compound of Formula I, II, III, IV, or V in combination with one or more. Formula I in the range of about 25 to 1000 mg once or twice daily and repeated doses of the anti-diabetic agent or agents at jointly effective dosages. The jointly effective dosage for PPAR modulators disclosed herein may be readily determined.

L12 ANSWER 28 OF 115 USPATFULL, on STN

2003:38352 USPATFULL ACCESSION NUMBER:

TITLE:

INVENTOR(S):

SUMM

DETD

143 human secreted proteins

Rosen, Craig A., Laytonsville, MD, UNITED STATES Ruben, Steven M., Olney, MD, UNITED STATES

Moore, Paul A., Germantown, MD, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Komatsoulis, George A., Silver Spring, MD, UNITED

STATES

Birse, Charles E., North Potomac, MD, UNITED STATES Duan, Roxanne D., Bethesda, MD, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES Soppet, Daniel R., Centreville, VA, UNITED STATES

NUMBER	KIND	DATE
	21'	20030206

PATENT INFORMATION: APPLICATION INFO .: RELATED APPLN. INFO.: US 2003027999 A1 20030206 20011108 (9) US 2001-986480 A1

Continuation-in-part of Ser. No. WO 2000-US12788, filed

on 11 May 2000, UNKNOWN DATE NUMBER 19990513 (60) US 1999-134068P PRIORITY INFORMATION: Utility DOCUMENT TYPE: APPLICATION HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, FILE SEGMENT: LEGAL REPRESENTATIVE: ROCKVILLE, MD, 20850 NUMBER OF CLAIMS: EXEMPLARY CLAIM: LINE COUNT: 29687 CAS INDEXING IS AVAILABLE FOR THIS PATENT. . . . component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders. . . . polypeptides corresponding to this gene and/or agonists and/or SUMM antagonists thereof may be used to diagnose, prognose, treat, prevent, and/or ameliorate type II diabetes mellitus (insulin resistant diabetes mellitus). . . neutralizing or antagonistic antibodies) may be used to SUMM diagnose, prognose, treat, prevent, and/or ameliorate conditions associated with (type I or type II) diabetes mellitus, including, but not limited to, diabetic ketoacidosis, diabetic coma, nonketotic hyperglycemichyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . renal failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and impaired wound healing, infections (e.g., infectious diseases and disorders as described in the "Infectious Diseases". (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. DETD (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. . . [2060] The diabetic animals have many of the characteristic DETD features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1992)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al., J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol. 83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67 (1980); Giacomelli et. al., Lab Invest. 40(4):460-473 (1979); Coleman, D. L., Diabetes 31 (Suppl): 1-6 (1982)). These homozygous

L12 ANSWER 29 OF 115 USPATFULL on STN
ACCESSION NUMBER: 2003:38129 USPATFULL
TITLE: 29 human cancer associated proteins

analogous to human type II diabetes

INVENTOR(S):

Z9 numan cancer associated proteins

Roschke, Viktor, Rockville, MD, UNITED STATES

(Mandel et al., J. Immunol. 120:1375-1377 (1978)).

diabetic mice develop hyperglycemia that is resistant to insulin

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DATE
                                          KIND
                             NUMBER
                                                20030206
                       US 2003027776
                                          A1
PATENT INFORMATION:
                                                         (10)
                                          A1
                                                20011221
                        US 2001-23896
                        Continuation-in-part of Ser. No. WO 2000-US23794, filed
APPLICATION INFO .:
RELATED APPLN. INFO.:
                        on 30 Aug 2000, UNKNOWN
                               NUMBER
                                           19990903 (60)
                        US 1999-152296P
PRIORITY INFORMATION:
                                           19991006 (60)
                        US 1999-158003P
                        Utility
DOCUMENT TYPE:
                        HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,
                        APPLICATION
FILE SEGMENT:
LEGAL REPRESENTATIVE:
                        ROCKVILLE, MD, 20850
NUMBER OF CLAIMS:
EXEMPLARY CLAIM:
                         23049
 LINE COUNT:
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
        . . . tissues and organs (e.g., cancers of the hypothalamus,
        pituitary gland, thyroid gland, parathyroid glands, pancreas, adrenal
        glands, ovaries, and testes), diabetes (e.g., diabetes
        insipidus, type I and type II diabetes
        mellitus), obesity, disorders related to pituitary glands (e.g.,
        hyperpituitarism, hypopituitarism, and pituitary dwarfism),
        hypothyroidism, hyperthyroidism, goiter, reproductive disorders (e.g.
              . . adrenal glands (e.g., Addison's Disease, corticosteroid
        deficiency, and Cushing's Syndrome), kidney cancer (e.g., hypernephroma,
        transitional cell cancer, and Wilm's tumor), diabetic
        nephropathy, interstitial nephritis, polycystic kidney disease,
        glomerulonephritis (e.g., IgM mesangial proliferative glomerulonephritis
        and glomerulonephritis caused by autoimmune disorders; such as. .
              . component that may be treated, prevented, and/or diagnosed with
         the compositions of the invention include, but are not limited to,
 SIJMM
         type II collagen-induced arthritis, antiphospholipid
         syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing
         polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia,
         polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune
         pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin
         dependent diabetes mellitus, and autoimmune inflammatory eye
           . . polypeptides corresponding to this gene and/or agonists and/or
         disorders.
         antagonists thereof may be used to diagnose, prognose, treat, prevent,
  SUMM
         and/or ameliorate type II diabetes
         mellitus (insulin resistant diabetes mellitus).
         . . . neutralizing or antagonistic antibodies) may be used to
         diagnose, prognose, treat, prevent, and/or ameliorate conditions
  SUMM
         associated with (type I or type II) diabetes
         mellitus, including, but not limited to, diabetic
         ketoacidosis, diabetic coma, nonketotic hyperglycemic
         -hyperosmolar coma, seizures, mental confusion, drowsiness,
         cardiovascular disease (e.g., heart disease, atherosclerosis,
         microvascular disease, hypertension, stroke, and. . . renal failure,
         nephropathy other diseases and disorders as described in the "Renal
         Disorders" section), nerve damage, neuropathy, vision impairment (e.g.,
         diabetic retinopathy and blindness), ulcers and impaired wound
         healing, infections (e.g., infectious diseases and disorders as
         described in the "Infectious Diseases".
                   (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM.
          (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose),
  DETD
          AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs)
          such as rosiglitazone, AVANDIA.TM. (rosiglitazone
          maleate) ACTOS.TM. (piogliatazone), and troglitazone;
          alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins
          such as SANDOSTATIN.TM. (octreotide); and diazoxides such as
          [1358] The diabetic animals have many of the characteristic
          PROGLYCEM. TM...
   DETD
          features observed in Type II diabetes
```

mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman et al. Proc. Natl. Acad. Sci. USA 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel et al., J. Immunol.. . . glomerular filtration abnormalities have been described in these animals (Norido, F. et al., Exp. Neurol. 83(2):221-232 (1984); Robertson et al., Diabetes 29(1):60-67 (1980); Giacomelli et al., Lab Invest. 40(4):460-473 (1979); Coleman, D. L., Diabetes 31 (Suppl): 1-6 (1982)). These homozygous diabetic mice develop hyperglycemia that is resistant to insulin analogous to human type II diabetes (Mandel et al., J. Immunol. 120:1375-1377 (1978)).

L12 ANSWER 30 OF 115 USPATFULL on STN

ACCESSION NUMBER:

2003:31119 USPATFULL

TITLE:

Attractin-like polynucleotides, polypeptides, and

antibodies

INVENTOR(S):

Ni, Jian, Germantown, MD, UNITED STATES Ruben, Steven M., Olney, MD, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Human Genome Sciences, Inc., Rockville, MD, UNITED

PATENT ASSIGNEE(S):

STATES, 20850 (U.S. corporation)

KIND NUMBER -----

PATENT INFORMATION:

US 2003023070 A1 20030130 US 2002-84994 A1 20020301 (10) RELATED APPLN. INFO.: Continuation of Ser. No. US 2001-790621, filed on 23 Feb 2001, PENDING Continuation-in-part of Ser. No. WO

2000-US23663, filed on 29 Aug 2000, UNKNOWN

NUMBER

DATE

PRIORITY INFORMATION:

US 1999-151348P 19990830 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

POCKVILLE, MD, 20850

NUMBER OF CLAIMS:

22 1

EXEMPLARY CLAIM:

LINE COUNT:

12029

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

SUMM . . . component that may be treated, prevented, and/or diagnosed with

the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, Neuritis, Uveitis Ophthalmia, Polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, Autoimmune

Pulmonary Inflammation, Autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye.

MMUR

. . . polypeptides corresponding to this gene and/or agonists and/or antagonists thereof may be used to diagnose, prognose, treat, prevent, and/or ameliorate type II diabetes

mellitus (insulin resistant diabetes mellitus).

SUMM

. . . neutralizing or antagonistic antibodies) may be used to diagnose, prognose, treat, prevent, or ameliorate conditions associated with (type I or type II) diabetes mellitus, including, but not limited to, diabetic

ketoacidosis, diabetic coma, nonketotic hyperglycemic-hyperosmolar coma, seizures, mental confusion, drowsiness, cardiovascular disease (e.g., heart disease, atherosclerosis, microvascular disease, hypertension, stroke, and other. . . failure, nephropathy other diseases and disorders as described in the "Renal Disorders" section), nerve damage, neuropathy, vision impairment (e.g., diabetic retinopathy and blindness), ulcers and

impaired wound healing, infections (e.g., infectious diseases and

DETD

disorders as described in the "Infectious Diseases". (glyburide), GLUCOTROL.TM. (glipizide), and DIAMICRON.TM. (gliclazide), GLUCOPHAGE.TM. (metformin), PRECOSE.TM. (acarbose), AMARYL.TM. (glimepiride), and ciglitazone; thiazolidinediones (TZDs) such as rosiglitazone, AVANDIA.TM. (rosiglitazone maleate) ACTOS.TM. (piogliatazone), and troglitazone; alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN.TM. (octreotide); and diazoxides such as PROGLYCEM.TM.. . .